Practice Parameters for the Surgical Treatment of Ulcerative Colitis

Howard Ross, M.D. • Scott R. Steele, M.D. • Mika Varma, M.D. • Sharon Dykes, M.D. Robert Cima, M.D. • W. Donald Buie, M.D. • Janice Rafferty, M.D.

Prepared by the Standards Practice Task Force of the American Society of Colon and Rectal Surgeons

he American Society of Colon and Rectal Surgeons is dedicated to assuring high-quality patient care by advancing the science, prevention, and management of disorders and diseases of the colon, rectum, and anus. The Standards Committee is composed of Society members who are chosen because they have demonstrated expertise in the specialty of colon and rectal surgery. This Committee was created to lead international efforts in defining quality care for conditions related to the colon, rectum, and anus. This is accompanied by developing Clinical Practice Guidelines based on the best available evidence. These guidelines are inclusive, and not prescriptive. Their purpose is to provide information on which decisions can be made, rather than dictate a specific form of treatment. These guidelines are intended for the use of all practitioners, health care workers, and patients who desire information about the management of the conditions addressed by the topics covered in these guidelines. It should be recognized that these guidelines should not be deemed inclusive of all proper methods of care or exclusive of methods of care reasonably directed to obtaining the same results. The ultimate judgment regarding the propriety of any specific procedure must be made by the physician in light of all of the circumstances presented by the individual patient.

STATEMENT OF THE PROBLEM

Ulcerative colitis (UC) is characterized by a chronic inflammatory condition that affects the rectum and extends proximally into the colon for varying distances. Although many patients are treated effectively by a wide variety of medications, approximately 15% to 30% of patients will require or elect operative intervention. Surgical indications vary from acute colitis to malignancy and often have a dramatic impact on morbidity, mortality, and quality of life.^{1,2} Failure to respond to medical therapy (ie, intractability) remains

Dis Colon Rectum 2014; 57: 5–22 DOI: 10.1097/DCR.00000000000000030 © The ASCRS 2013 the most common surgical indication for UC. Whether it is secondary to an inability to control symptoms, poor quality of life, risks/side effects of chronic medical therapy (especially long-term corticosteroids), noncompliance, or growth failure, patients may opt for surgery in the elective or semielective setting.³ Complete removal of all the potential disease-bearing tissue is theoretically curative in UC. Operative options include an abdominal colectomy or total proctocolectomy with either a permanent end ileostomy or surgical construction of a "new" rectum through an IPAA that restores GI continuity.^{4,5} All these procedures may be performed by using open or minimally invasive techniques.^{6,7} Although various treatment options are available for UC, this parameter will focus on the surgical management for the patient with UC.

METHODOLOGY

These guidelines are built on the last set of the American Society of Colon and Rectal Surgeons Practice Parameters for treatment of UC published in 2005.8 An organized search of Medline, PubMed, and the Cochrane Database of Collected Reviews was performed through July 2013. Keyword combinations included inflammatory bowel disease, ulcerative colitis, ileal pouch-anal anastomosis, ileostomy, proctocolectomy, colorectal neoplasm, surgery, colectomy, ileoproctostomy, immunomodulator, infliximab, steroids, and related articles. Directed searches of the embedded references from the primary articles also were accomplished. The final grade of recommendation was performed using the Grades of Recommendation, Assessment, Development, and Evaluation (GRADE) system (Table 1).9

INDICATIONS FOR SURGERY

Acute Colitis

1. Patients with clinical evidence of actual or impending perforation should undergo urgent surgery. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

TABLE 1. The GRADE system-grading recommendations				
	Description	Benefit vs risk and burdens	Methodological quality of supporting evidence	Implications
1A	Strong recommendation, high- quality evidence	Benefits clearly outweigh risk and burdens or vice versa	RCTs without important limitations or overwhelming evidence from observational studies	Strong recommendation, can apply to most patients in most circumstances without reservation
1B	Strong recommendation, moderate-quality evidence	Benefits clearly outweigh risk and burdens or vice versa	RCTs with important limitations (inconsistent results, methodological flaws, indirect or imprecise) or exceptionally strong evidence from observational studies	Strong recommendation, can apply to most patients in most circumstances without reservation
1C	Strong recommendation, low- or very-low- quality evidence	Benefits clearly outweigh risk and burdens or vice versa	Observational studies or case series	Strong recommendation but may change when higher-quality evidence becomes available
2A	Weak recommendation, high- quality evidence	Benefits closely balanced with risks and burdens	RCTs without important limitations or overwhelming evidence from observational studies	Weak recommendation, best action may differ depending on circumstances or patients' or societal values
2B	Weak recommendations, moderate-quality evidence	Benefits closely balanced with risks and burdens	RCTs with important limitations (inconsistent results, methodological flaws, indirect or imprecise) or exceptionally strong evidence from observational studies	Weak recommendation, best action may differ depending on circumstances or patients' or societal values
2C	Weak recommendation, low- or very-low- quality evidence	Uncertainty in the estimates of benefits, risks and burden; benefits, risk and burden may be closely balanced	Observational studies or case series	Very weak recommendations; other alternatives may be equally reasonable

GRADE = Grades of Recommendation, Assessment, Development, and Evaluation; RCT = randomized controlled trial.

Adapted from Guyatt G, Gutermen D, Baumann MH, et al. Grading strength of recommendations and quality of evidence in clinical guidelines: report from an American College of Chest Physicians Task Force. Chest. 2006;129:174–181. Used with permission.

The diagnosis of *severe* colitis is based on the criteria of Truelove and Witts¹⁰ and is defined as colitis with more than 6 bloody stools per day, fever (temperature, >37.5°C), tachycardia (heart rate, >90 beats per minute), anemia (hemoglobin, <75% of normal), and elevated sedimentation rate (>30 mm/h).¹¹ *Toxic or fulminant* colitis is characterized by more than 10 bloody stools per day, fever (>37.5°C), tachycardia (>90 beats per minute), anemia requiring transfusion, elevated sedimentation rate (>30 mm), colonic dilation on radiography, and abdominal distention with tenderness.¹¹ When the colonic distention of the transverse colon exceeds 6 cm, the diagnosis becomes *toxic megacolon*.^{12,13} Surgery is required in 20% to 30% of patients with toxic colitis, and typically consists of a subtotal colectomy with end ileostomy. ^{14,15}

Perforation in patients with toxic colitis is associated with a high mortality rate (27%–57%), regardless of whether the perforation is contained or free. The mortality rate also increases as the time interval between perforation and surgery increases. Signs of impending perforation may be masked by ongoing medical therapy. Persistent or increasing colonic dilation, pneumatosis coli, worsening local peritonitis, and the development of multiple organ failure can be signs of impending or ac-

tual perforation.^{17,19,20} Likewise, localized peritonitis may reflect local inflammation or may be a sign of impending perforation.²¹ Perforation may also occur without dilation; these patients often do not exhibit classic signs of peritonitis.¹⁶

The development of multisystem organ failure (MSOF) is an ominous sign. In a series of 180 patients with toxic colitis, 11 (6%) developed MSOF. The overall mortality in the entire group was 6.7%; however, of the 12 patient deaths, 8 occurred in patients with MSOF.²²

2. For patients with moderate to severe colitis, early surgical consultation should be obtained. Grade of Recommendation: Strong recommendation based on low-quality evidence, 1C.

Although most physicians agree that an urgent surgical consultation should be obtained in patients with sepsis or fulminant/toxic colitis, there is no high-quality literature depicting the appropriate timing for surgical consultation in patients with a lesser degree of disease. Recent consensus statements from surgical and gastroenterology experts have suggested that a failure of primary therapy or those patients being considered for monoclonal antibody or cyclosporine therapy warrant surgical consultation.²³ It

is important to keep in mind that, even in those patients who have a good initial response to medical therapy, the eventual need for colectomy ranges from 20% to 80%. 24-27 Early involvement of the surgeon is important to not only follow the patient's clinical course, but also to provide information and answer questions should the need for surgical therapy eventually arise. This is best performed in an elective setting rather than urgently in a patient who has perforated or is clinically deteriorating. Concomitant involvement of an enterostomal therapist is also valuable in providing education and perioperative ostomy marking should the need arise. ^{28,29}

3. Patients whose condition worsens on medical therapy or who do not make significant improvement after a period of 48 to 96 hours of appropriate medical therapy should be considered for either a second-line agent or surgery. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Medical therapy is considered to have failed in patients if their condition worsens while on medical therapy, or if they do not improve after a period of initial stabilization. Monoclonal antibody therapy (eg, infliximab, adalimumab) has demonstrated effectiveness for rescue therapy for hospitalized patients.^{30–32} The short-term response rate, defined as avoidance of colectomy and cessation of corticosteroids, is ~25% (range, 20%-90%). The ability to avoid colectomy following an acute episode in the short term for patients refractory to steroids has been reported in up to 90%,³³ although the impact of infliximab for rescue therapy is still evolving. Concerns about using biological agents are related to the large percentage of patients who ultimately require colectomy, and safety issues that are primarily infectious complications.^{33–38} Limited evidence suggests that intravenous cyclosporine is more effective than standard steroid-based treatment for severe colitis39-41 and has been advocated as a second-line agent before colectomy, in part, secondary to its rapid onset of action and short half-life.42 Sequential "rescue" therapy with either monoclonal antibody therapy or cyclosporine following failure of the other has been associated with poor outcomes and is generally not recommended.^{43–45}

The need for and timing of surgery in patients whose condition seems to "plateau" after a period of initial improvement often is difficult to judge. Patients with more than 8 stools per day, or 3 to 8 stools and a C-reactive protein > 45 mg/mL after 3 days of therapy, have an 85% chance of requiring colectomy during the same hospitalization, regardless of whether corticosteroid or cyclosporine treatment is used. 46 Patients who have a contraindication to (or do not desire) monoclonal antibody or cyclosporine therapy, or when steroids fail, should be considered for surgery. Furthermore, persistent colonic distention seems to characterize a subgroup of patients who respond

poorly to medical therapy and are at increased risk for the development of megacolon.⁴⁷ Prolonged observation of these patients may risk exhaustion of their physiological reserve, but does not necessarily increase perioperative morbidity outside of a perforation.¹⁵ Most series define a period of 48 to 96 hours after which surgery is indicated if the patient does not improve or worsens,^{14,18,19} although evidence specifying the most appropriate time period for a trial of medical therapy, especially with "second-line" agents, is lacking.

4. A decision regarding the response to second-line or "rescue" therapy should be made within 5 to 7 days after initiation. Grade of Recommendation: Strong recommendation based on low-quality evidence, 1C.

Both cyclosporine and monoclonal antibody therapy have been shown to have mean response times of 5 to 7 days in controlled trials.^{31,39,40} Population-based data evaluating timing of colectomy in UC have also demonstrated that mortality rates increase as operative timing progresses from within 3 to 6 days (OR, 2.12; 95% CI, 1.13–3.97) and 11 days (OR, 2.89; 95% CI, 1.41–5.91).⁴⁸ Longer waiting times may result in worsening physiological reserve, further depletion in nutrition stores, and an inappropriate delay in surgery with no apparent gain.⁴⁹

Intractability

1. Surgery is indicated in chronic UC when medical therapy is ineffective. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Intractability is one of the most common surgical indications for UC. Symptoms may be insufficiently controlled, despite an intensive medical regimen, and the patient is unable to achieve an acceptable quality of life. Alternatively, the response to treatment may be adequate, but the risks of chronic medical therapy may be excessive. Additionally, patients who are unable to tolerate the deleterious side effects of medical therapy and those patients who are noncompliant with treatment regimens may be candidates for surgical management.

Disabling extraintestinal manifestations of UC may prompt resection. Typically, episcleritis, erythema nodosum, aphthous ulcerations, and large joint arthropathy are more likely to be responsive to colectomy. Hepatic, vascular, hematologic, cardiopulmonary, and neurological comorbidities typically are not. Finally, growth failure in children is another form of intractability that may require colectomy. Surgery should be considered if growth failure persists despite maximal nutritional and medical therapy, with data reporting that surgical resection is at least as effective as immunosuppressants for allowing "catch-up" in growth status. ^{50–52}

Cancer Risk and Surveillance

1. Patients with long-standing UC should undergo endoscopic surveillance. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Surveillance recommendations in UC are based on patient risk factors, current literature, and expert panel consensus with the goal of early detection of premalignant and malignant lesions, along with decreasing mortality. Risk factors associated with the development of malignancy include the extent of colonic disease, pancolitis (proximal to the splenic flexure) greater than left-sided colitis, and prolonged disease duration (>8 years with pancolitis).^{53,54} Consensus evidence also remains that a diagnosis of UC at a younger age is associated with an increased risk of colorectal cancer.53 Additional risk factors include family history of IBD55 and concomitant primary sclerosing cholangitis.56-58 Although a meta-analysis demonstrated the cumulative risk of colorectal cancer for UC patients to be 2.1% at 10 years, 8.5% at 20 years, and 17.8% at 30 years,⁵⁹ population-based series have reported lower annual incidence rates of 0.06% to 0.2%. 60,61 Despite this, it is generally accepted that chronic UC is associated with an increased risk of malignancy.

Surveillance colonoscopy is recommended for these patients, even though data supporting a survival benefit are lacking. A Cochrane update reported that there is "no clear evidence" that surveillance colonoscopy is associated with longer survival in patients with extensive colitis (relative risk, 0.81; 95% CI, 0.17–3.83).⁶² The authors did note that cancers are generally detected at an earlier stage in those undergoing regular surveillance endoscopy, with a corresponding associated overall better prognosis.

Patients with extensive colitis (disease proximal to the splenic flexure) should be advised to undergo endoscopy after 8 years of disease and should have a surveillance colonoscopy performed every 1 to 2 years. This interval will depend on the presence or absence of dysplasia in biopsy specimens (see below). Previous recommendations based on expert opinion suggest that, for patients with left-sided disease (disease distal to the splenic flexure yet proximal to the rectum), the same surveillance programs may begin after 15 years of the disease; 63-66 however, recent American Gastroenterological Association guidelines suggest that the evidence is not strong enough to recommend different surveillance intervals.⁶⁷ Patients with 2 successive negative surveillance colonoscopies may undergo surveillance colonoscopy at 1 to 3 years, although this should be tailored to the patient. 63 There are additional data to indicate that patients with concomitant primary sclerosing cholangitis are at an even higher risk for malignancy, with a cumulative risk of cancer or dysplasia approaching 50% at 25 years of disease.⁵⁸ Therefore, it is recommended that these patients adhere to annual surveillance endoscopy regardless of previous normal findings.

2. Endoscopic surveillance should involve 2 sets of 4-quadrant random biopsies at ~10-cm intervals throughout the colon and rectum, along with directed biopsies of suspicious lesions. Grade of Recommendation: Strong recommendation based on low-quality evidence, 1C.

Recommendations vary slightly based on society and expert opinion. ^{66–69} The recommended surveillance endoscopy technique normally involves 2 sets of 4-quadrant biopsies in each colon segment (right, transverse, left, rectosigmoid), producing ~32 random biopsies divided into 4 specimen cups. A minimum of 32 random biopsies has been shown to result in an 80% to 90% sensitivity for detecting dysplasia. ^{70,71} Directed biopsies of polypoid lesions, masses, strictures, or irregular mucosa distinct from surrounding inflammation should also be performed. ^{72,73} Additionally, adjacent "normal"-appearing tissue, when present, should be sent for comparison. Polyps that appear potentially dysplastic can be removed by polypectomy, and the adjacent flat mucosa also should be biopsied to exclude dysplasia.

In an attempt to aid in identifying more subtle, flat, dysplastic lesions, targeted biopsies with the use of magnification chromoendoscopy have been described. In this technique, a stain/dye such as indigo carmine or methylene blue is applied throughout the entire colon and rectum to enhance mucosal changes that are less well visualized by standard techniques.74 This may be combined with advanced imaging capabilities including narrow-band imaging and confocal laser endomicroscopy to further augment visualization of dysplastic areas.⁷⁵ A small number of prospective studies have demonstrated significant improvement in intraepithelial neoplasia detection over conventional light endoscopy.^{76–78} Although this promising development may be considered, further widespread availability and collective experience outside of select centers is required before generalized recommendations.

3. Total proctocolectomy, with or without IPAA, is recommended for patients with carcinoma, non-adenomalike dysplasia-associated lesion or mass, or high-grade dysplasia. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Malignancy in UC patients is felt to be secondary to similar genetic mutations responsible for sporadic colon cancer in the general population, as well as persistent inflammation resulting in cell proliferation, oxidative stress, and, ultimately, dysplasia.^{79,80} Unlike sporadic cases, cancer in UC patients may not always follow a progression from normal epithelium to low-grade dysplasia, high-grade dysplasia, and finally invasive malignancy. Nevertheless, dysplasia detection by conventional histopathologic assessment of colonoscopic biopsies remains the criterion standard to identify patients at highest risk of developing cancer in UC.⁸¹ Consideration should be made for all biopsies con-

cerning for high-grade dysplasia to be confirmed by 2 independent GI pathologists when possible.

Findings of colorectal cancer, a *nonadenoma* dysplasia-associated lesion or mass (DALM), or high-grade dysplasia are almost uniformly accepted as indications for proctocolectomy. With or without IPAA, because approximately 43% to 50% will have concomitant malignancy at the time of colectomy. Page 10 a classic study evaluating 10 prospective cohorts including 1225 patients undergoing surveillance colonoscopy, 10 of 24 patients (42%) found to have high-grade dysplasia who underwent immediate colectomy were found to have a synchronous cancer. Page 20 and 20

Adenoma-like DALM may occur in areas outside of colitis and may be successfully treated via endoscopic polypectomy similar to sporadic adenomas. Recent studies suggest that adenoma-like DALM located within regions of active inflammation may also be removed endoscopically with the avoidance of colectomy, given a complete resection and no evidence of adjacent flat dysplasia or adenocarcinoma. However, when the polypectomy base or surrounding mucosa does contain dysplasia, patients should undergo surgical resection.

4. Total proctocolectomy, or surveillance endoscopy, is recommended for patients with UC and low-grade dysplasia. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1C.

The management of unifocal, flat, low-grade dysplasia in the setting of UC remains controversial. Reported progression rates to high-grade dysplasia range widely from 0% to 53%. 92-95 A recent meta-analysis including 20 colonoscopic surveillance studies and 508 patients with flat low-grade dysplasia with DALM, reported a 9-fold risk of developing cancer (OR, 9.0; 95% CI, 4–20.5) and 12-fold risk of developing any advanced lesion (OR, 11.9; 95% CI, 5.2–27). 96 Bernstein et al 22 confirmed the risk of cancer in the setting of low-grade dysplasia as 19%; however, in up to 29% of the patients they followed, untreated low-grade dysplasia went on to develop a DALM, high-grade dysplasia, or cancer, prompting the authors to recommend immediate colectomy for these patients.

In a conflicting study, 60 patients with low-grade dysplasia in flat mucosa identified during endoscopy were followed for an average of 10 years; low-grade dysplasia was seen at several locations and, during repeated colonoscopies, in 73% of cases. However, progression to high-grade dysplasia or a dysplasia-associated lesion/mass occurred in only 11 patients (18%). Pathough high rates of interobserver variation between histopathologists confound the findings and recommendations, 14,97–99 it is important that patients are counseled about the potential risks and benefits of continued endoscopic surveillance versus surgical therapy.

Patients may benefit from chemoprevention with 5-aminosalicylate (5-ASA), but there are few prospective

data evaluating this topic. A recent meta-analysis of 9 observational studies reported a reduced risk of developing colorectal carcinoma or dysplasia with 5-ASA use (OR, 0.51; 95% CI, 0.38–0.69). Case-control series have demonstrated a significant risk reduction (OR range, 0.19–0.60, all p < 0.05) for both dysplasia and colorectal cancer in patients using 5-ASA compounds. Other large population-based and case-control studies have not found a protective effect with regular 5-ASA exposure. Nevertheless, the side effects of these medications should be compared with the potential for reducing malignancy when counseling patients.

5. Patients with UC who develop a stricture, especially with long-standing disease, should typically undergo resection. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Among the most common manifestations of colorectal carcinoma in chronic UC are colonic strictures, which develop in 5% to 10% of patients with UC. Although the majority of strictures are benign, as many as 25% will be malignant, and malignant strictures account for up to 30% of cancers occurring in UC patients. Although biopsy may reveal dysplasia or malignancy, ¹⁰⁶ a negative biopsy may not be reliable because of the risk of sampling error and the more infiltrative nature of colitis-associated malignancies. Therefore, in general, all patients with strictures should undergo an oncological resection. ^{107,108}

SURGICAL OPTIONS

Emergency

1. The procedure of choice for emergency surgery in UC is total or subtotal abdominal colectomy with end ileostomy. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

The surgical goals in the acute setting are designed to remove the bulk of the diseased bowel, restore patient health with the greatest reliability and least risk, and preserve reconstructive options after the patient has recovered and medications are withdrawn.⁵ Subtotal colectomy with end ileostomy and Hartmann closure of the distal bowel or creation of a mucous fistula is a safe and effective approach.^{15,109,110} Although, historically, this has been performed via a laparotomy, multiple reports have confirmed the feasibility and safety of a minimally invasive approach in this setting.111-113 Extrafascial placement of a closed rectosigmoid stump may be associated with fewer pelvic septic complications and facilitates subsequent pelvic dissection.114 Transanal drainage of the distal stump may further decrease the risk of pelvic sepsis.¹¹⁵ The resected colon specimen should be examined microscopically for confirmation of UC or Crohn's disease, because the likelihood of an altered diagnosis is appreciable after colectomy. ^{15,113} In patients with confirmed UC, a completion proctectomy and IPAA can often be safely performed at a later date to remove the remaining disease and restore intestinal continuity. Although there are reports of successful outcomes following total proctocolectomy with IPAA in select patients with a "moderate" form of fulminant colitis, in general, this should be avoided. ^{116,117}

Elective Surgery

1. Total proctocolectomy with ileostomy is an acceptable surgical option for patients with UC. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Total proctocolectomy with ileostomy has been the conventional operative approach for patients with UC and may be considered a benchmark procedure with which all other operations are compared.^{6,7} It is a safe, effective, and curative operation that permits most patients to live a full, active lifestyle.¹¹⁸ Although restorative proctocolectomy with IPAA has become increasingly popular during the past 3 decades, proctocolectomy with ileostomy can still be considered the first-line procedure for patients who choose not to undergo a restorative proctocolectomy and for those at significant risk for pouch failure, such as patients with impaired anal sphincter muscles, previous anoperineal disease, or limited physiological reserve secondary to comorbid conditions.^{119,120} When the expertise is available, minimally invasive surgery can be considered.¹²¹

Regardless of technique, the operation does have recognized complications. Although stoma-associated problems, such as stenosis and prolapse, are probably the most frequent, 122 other complications that are common to any abdominal/pelvic procedure also have been recognized following this operation. 122,123 These include small-bowel obstruction, infection/fistula, persistent pain, unhealed perineal wound, sexual and bladder dysfunction, and infertility. 124,125 In 1 study of 44 patients, the long-term complication rate of proctocolectomy with permanent ileostomy was significantly lower than restorative proctocolectomy (26% vs 52%). 123 There have also been several recent case reports of carcinoma arising at the ileostomy site, similar to that of the rectal stump, highlighting the need for continued surveillance. 126–129

2. Total proctocolectomy with IPAA is an appropriate operation for selected patients with UC. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Total proctocolectomy with IPAA has become the most commonly performed procedure for patients with UC who require elective surgery. The operation, whether performed through an open procedure or, when expertise is available, by the use of minimally invasive techniques, is relatively safe and durable. ^{130–135} Ileal pouch-anal anastomosis is associated with an acceptable morbidity rate (19%–27%), ^{136,137} an extremely low mortality rate (0.2%–0.4%), ^{136–142} and a quality of life that approaches that of the healthy population. ^{143–153} When deciding on IPAA, patient selection should consider factors such baseline continence, ability to undergo major pelvic surgery and its complications, and medical comorbidities.

The complications of the procedure are similar to those of any major abdominal operation, but they also include risks arising from the pelvic dissection, such as infertility or sexual dysfunction, and pouch-specific complications, such as pouchitis. Additionally, anastomotic leak with pelvic sepsis, fistula, stricture, and cuff inflammation may occur. Medical therapies, such as antibiotics, immunomodulators, and biologics, as well as surgery, may be attempted to salvage pouch function and are successful in more than 50% of cases. ^{154–156}

When comparing the open versus a laparoscopic approach, a 2009 Cochrane review consisting of 11 trials and more than 600 patients found similar length of stay, morbidity, reoperation, readmission, and mortality rates between the 2 approaches.¹⁵⁷ Laparoscopy was associated with improved cosmesis, smaller total incision length, and longer operative times. Comparing 2-stage procedures, retrospective data have demonstrated that laparoscopy is associated with a shorter time to ileostomy closure and restoration of intestinal continuity, likely secondary to decreased adhesion formation.¹⁵⁸ Studies using diagnostic laparoscopy at the time of ileostomy closure have confirmed lower adhesion formation with a minimally invasive approach, whether in the incisional, abdominal, or adnexal location, 159 including >70% with no adnexal adhesions.160

The impact of moderate- to high-dose steroids on the incidence of postoperative complications after IPAA is well described. However, the impact of infliximab in this setting is less clear. There is a larger experience with these agents and surgery for Crohn's disease that has not clearly demonstrated a relationship between α -tumor necrosis factor agents and postoperative complications. 162 Unfortunately, the literature on the impact of these agents in patients with chronic UC undergoing IPAA is limited to retrospective single-institution studies with varying patient numbers with differing procedure mixes and variable definitions of complications. $^{143,163-168}$

Currently, reports regarding the impact of α -tumor necrosis factor therapy before IPAA on postoperative complications are limited to observational studies, have different patient populations, and lack a consistent definition of complications. Further larger studies, ideally multi-institutional, using standardized procedures and complication definitions will be required to identify any

association with the preoperative use of these agents and poor outcomes after IPAA.

3. Patients with UC considering pelvic operations should be counseled regarding the potential negative effects on sexual function and fertility. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

Although many studies have shown no difference in fertility in women with UC compared with healthy controls, 169,170 those who undergo surgical treatment for UC have a lower rate of conception compared with their nonsurgical counterparts. 144-147,170-172 The results of a metaanalysis indicate a 3-fold increase in infertility in women with UC undergoing surgical therapy, particularly IPAA, compared with either healthy controls or those with UC who do not have surgery.¹⁷² Overall, infertility rates after IPAA range from 38% to 64% compared with only 10% to 38% for nonoperative patients or controls. 144-147,170-172 Furthermore, a greater percentage of women after IPAA require fertility treatments (18% vs 6%). 145,170 Pelvic adhesions and occlusion of the fallopian tubes are the proposed reasons for this difference. This seems to be associated with the rectal resection component and not pouch placement itself, because ileorectal anastomosis in UC has not been associated with decreased fertility.¹⁷³

Rates of postoperative fertility problems are similar between proctocolectomy with IPAA and with end ileostomy following other types of rectal resection such as familial adenomatous polyposis. 174 Yet, in the UC population, the literature has focused more on fertility following IPAA than after total proctocolectomy with end ileostomy. Multicenter data have demonstrated that a minimally invasive approach is associated with significantly lower rates of infertility in comparison with traditional open surgery, citing conception rates of 31% to 73% following a straight laparoscopic IPAA. 175,176 Pregnancy after IPAA is also not associated with an increase in maternal or fetal complications, 144,177 although an increase in stool frequency and pad usage has been reported in the third trimester.¹⁴⁴ Vaginal delivery appears to be safe after IPAA, although this is based on small retrospective series. 144,177-179 Anterior sphincter defects have been reported in up to 50% of women undergoing vaginal delivery, although this does not seem to significantly influence pouch function.^{180,181} After delivery, transient anal dysfunction normally resolves^{177,182} and pouch function returns to pregestational function within 6 months,144 independent of the mode of delivery.183

Sexual function for both men and women after proctectomy for UC has been variable following surgery, with some reports demonstrating improvement and others deterioration. Several early studies in women noted increased frequency of sexual function, with normal sexual desire, ability to achieve orgasm and minimal sexual dissatisfaction; however, these studies were retrospective and

did not use validated measures. ^{184–186} There was a clear increase in dyspareunia after surgery ranging from 11% to 22%. ^{144,185,186} Rates of retrograde ejaculation and erectile dysfunction in men range from 2% to 19% and 0% to 15%. ^{185–187} The largest retrospective review of sexual function in 1454 men and women after IPAA reported no change in 56%, better function in 25%, and worse function in 19%. ¹⁸⁸ More recent studies using validated measures ^{189,190} have demonstrated that, although men may have improved sexual function, women may have no improvement or even worsening sexual function, especially with IPAA at an early age. ¹⁹¹ Further prospective studies are needed to confirm these findings.

4. Total proctocolectomy with IPAA may be offered to selected UC patients with concomitant colorectal cancer. Grade of Recommendation: Strong recommendation based on low-quality evidence, 1C.

Ulcerative colitis patients with a concomitant carcinoma have postoperative complications and functional results comparable to colitis patients without cancer. ^{192–198} Nearly 20% of UC patients with cancer who underwent an IPAA subsequently die of metastatic disease. ¹⁹⁷ A more conservative management approach has been advocated by some who recommend an abdominal colectomy with ileostomy followed by a restorative proctectomy after an observation period of at least 12 months to better ensure that no recurrent or distant disease develops. ¹⁹⁸

Metastatic disease is generally considered a contraindication to IPAA. These patients are ideally managed with segmental colectomy or abdominal colectomy with anastomosis to facilitate early discharge and allow them to pursue chemotherapy and spend the rest of their lives relatively free of complications relating to extensive surgery. Another group of patients who may not be eligible for IPAA are those with cancer of the mid or low rectum. Performing a standard oncological resection with total mesorectal excision, ensuring adequate margins and sphincter preservation, may allow IPAA construction after completion of adjuvant therapy. Importantly, adjuvant radiotherapy, if indicated, should be performed *preoperatively* whenever possible, because postoperative radiotherapy is associated with a high incidence of pouch loss secondary to radiation enteritis and poor pouch function. 192,194 Ulcerative colitis patients with cecal cancers represent another unique subgroup of patients. If a long segment of adjacent distal ileum with its mesenteric vessels must be sacrificed, difficulties with positioning of the reservoir into the pelvis may ensue, and an ileostomy may be necessary if a tension-free anastomosis cannot be attained.

5. Total proctocolectomy with IPAA may be offered to selected elderly patients with UC. Grade of Recommendation: Strong recommendation based on low-quality evidence, 1C.

Ileal pouch-anal anastomosis in select elderly patients is safe and feasible. 199–203 Chronologic age should not by itself be used as an exclusion criterion. However, careful consideration should be given to underlying comorbidities, as well as the patient's mental status and anal sphincter function. Pouch procedures are feasible in suitably motivated elderly individuals who understand the risks and potential functional difficulties that often accompany this procedure. Although some series have found that bowel frequency remains constant in the first decade after the surgical procedure, 204 others have found that the number of daytime and nighttime stools increases, as does the likelihood of fecal incontinence. 130

Mucosectomy and double-stapled procedures are both acceptable techniques in most circumstances. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

The potential advantages of the double-stapled approach include enhanced technical ease, because this approach avoids mucosectomy and the perineal phase of the operation, less tension on the anastomotic suture line, and possibly improved functional results.²⁰⁵ Sphincter injury is minimized²⁰⁶ and the anal transition zone, with its abundant supply of sensory nerve endings, is preserved. Three prospective, randomized trials and 1 comparative study have demonstrated no significant difference in perioperative complications or functional results for patients in whom a mucosectomy was performed versus those patients in whom the proximal anal canal mucosa was preserved with a double-stapled procedure. 207-210 Whereas the majority of patients undergo a double-stapled procedure, it is important that the surgeon performing an IPAA be familiar with both techniques in the event of failure or the inability to use a surgical stapler.²¹¹

7. Pouch configuration may be chosen based on individual surgeon preference. Grade of Recommendation: Weak recommendation based on moderate-quality evidence, 2B.

Several ileal pouch configurations have been devised in an attempt to reduce pouch complications and improve functional outcome. These include the double-loop J-pouch, the lateral isoperistaltic H-pouch, the triple-loop S-pouch, and the quadruple-loop W-pouch.²¹²⁻²¹⁴ Two randomized trials comparing the J-pouch and W-pouch did not substantiate an improvement in functional outcomes with a larger reservoir. 215,216 In 1 study, the median number of stools per day was the same in patients with a J-pouch or a W-pouch, and there was no difference between the 2 reservoirs in the rates of incontinence, urgency, soiling, and the use of antidiarrheal agents. 215 An S-pouch can provide additional length (2-4cm) compared with the J-pouch, and may help to minimize anastomotic tension.²¹⁷ However, the 2-cm exit conduit of the S-pouch may elongate with time, and obstructed defecation may develop.

8. In carefully selected patients, a 1-stage IPAA can be considered. Grade of Recommendation: Strong recommendation based on low-quality evidence, 1C.

Numerous centers have published series supporting the omission of a protective ileostomy at the time of restorative proctocolectomy in patients who meet certain criteria.^{218–223} Although they report similar anastomotic leak rates and pelvic sepsis-related complications for both diverted and nondiverted groups, in all of these studies, the use of protective ileostomy was left to the discretion of the surgeon—a potential source of bias. In general, those patients undergoing 1-stage procedures have been younger, healthier, less obese, without anemia or hypoalbuminemia, and on either no immunosuppressive medications or lower dosages.²²⁴ Furthermore, all technical aspects of the surgery were straightforward, with no excess blood loss, good blood supply to the ileal pouch, no anastomotic tension, and a visibly intact anastomosis. A few studies have strongly encouraged the abandonment of this practice owing to higher anastomotic leak rates, more life-threatening septic complications, and the need for subsequent emergent operation. 225-227 The nondiverted patients also appear to have a higher incidence of postoperative ileus, although the overall number of hospital days is less than for those with an ileostomy. 223,228,229 Although it is clear that the use of a protective ileostomy does not alter the incidence of anastomotic leak, the milder pelvic sepsis related to fecal diversion may have a favorable impact on subsequent pouch function or retention.²³⁰ This advantage must be weighed against the significant complications associated with both creation and takedown the ileostomy, including dehydration, wound infection, ileostomy closure leak and fistula, ileoanal anastomotic stricture, and small-bowel obstruction. 220,223,228,229,231-234 As in any operation, proper patient counseling regarding risks and benefits, along with informed consent, are imperative.

 Continent ileostomy is an alternative for patients with UC who are not eligible for or have had a failed restorative proctocolectomy. Grade of Recommendation: Weak recommendation based on moderate-quality evidence, 2B.

The present role of the continent ileostomy, also known as the Kock pouch, is primarily confined to patients with poor sphincter function or a failed IPAA, or to those who are dissatisfied with a conventional Brooke ileostomy.^{235,236} This reduced role is the result of the success of the IPAA and the high rate of early and late complications associated with the continent ileostomy.²³⁷

Early complications, most commonly sepsis (secondary to suture line leaks, fistulas, and stomal necrosis) and obstruction, are seen in approximately one-fourth to one-third of patients. ^{235,238,239} Late complications occur in up to 60% of patients and include incontinence and ob-

struction secondary to disruption or dysfunction of the valve.²³⁵ Valve revision is required in up to 60% of patients.²³⁷ Although valve prolapse has been reduced with stapling techniques,^{240,241} the overall pouch failure rate has not decreased.²⁴²

Initial pouch retention rates have been reported in up to 96% after 1 year, ^{235, 237} with a 71% cumulative success rate at 29 years in 1 study.²³⁷ The failure rate is greater after a previously failed IPAA (46%) than after primary construction (23%),²⁴² as well as being higher after conversion from conventional ileostomy than when created primarily.²⁴³ For the two-thirds of patients with a functional continent ileostomy, the reported quality of life is similar to that described for patients with IPAA.^{235,237,244} Largely, this procedure has been relegated to surgeons with the expertise and experience of not only pouch construction, but also the aftercare of these patients as well.

10. Total abdominal colectomy with ileoproctostomy may be considered only in a highly selected group of patients with UC. Grade of Recommendation: Weak recommendation based on moderate-quality evidence, 2B.

Total abdominal colectomy with ileoproctostomy requires a relatively normal rectum to create a safe anastomosis; severe rectal inflammation and marked decrease in rectal compliance are contraindications to the procedure. ^{245,246} Severe anoperineal disease, although unusual in UC, also precludes an ileorectal anastomosis. ²⁴⁷ Caution should be exercised with this procedure in the setting of colonic dysplasia, or carcinoma in a potentially curative situation. ²⁴⁸

The benefits of total abdominal colectomy with ileoproctostomy are its relative simplicity and predictability compared with IPAA. Preliminary results from a recent study also suggest that ileorectal anastomosis preserves female fertility.²⁴⁹ The disadvantages are related to the long-term durability of the procedure. Studies demonstrate a 12% to 53% failure rate with follow-up of at least 3.5 years.^{250–253} In addition, the theoretical risk of developing cancer in the remaining rectum should be considered when counseling the patient about surgical options. Although the incidence of developing cancer seems low (0%–8% with long-term follow-up),^{253–256} patients undergoing total abdominal colectomy with ileorectal anastomosis must be willing to undergo annual endoscopic screening.^{246–248,251–254}

POSTOPERATIVE CONSIDERATIONS

1. Routine surveillance of ileal pouches for dysplasia in the ileal mucosa is not warranted. Grade of Recommendation: Strong recommendation based on low-quality evidence, 1C.

Dysplastic and neoplastic transformation within the pouch performed in UC seems to be extremely rare, unlike the low, although slightly higher, rate seen in familial ade-

nomatous polyposis.^{257–263} A decrease in villous height and increase in concentration of crypts have been observed in most ileal pouches.²⁶⁴ These metaplastic changes of the ileal mucosa to a colonic type mucosa are considered adaptations to the reservoir function of the pouch. Outside of symptoms, routine surveillance of the pouch does not appear to be beneficial or warranted.

2. Surveillance of the residual rectal cuff or the anal transition zone following restorative proctocolectomy may detect malignant degeneration. Grade of Recommendation: Strong recommendation based on weak-quality evidence, 1C.

Although little evidence supports routine surveillance of the ileal pouch itself, ^{257,265} short-segment inflammation (ie, "cuffitis") ^{266–268} and perianastomotic zone carcinoma ^{269–273} are infrequent, but more legitimate concerns. Rare instances of adenocarcinoma in the pouch outlet, or anal transition zone, have been reported following both mucosectomy and stapled anastomoses—in some cases, more than 10 years following the initial operation. ^{274–276} Although the optimal surveillance interval remains largely anecdotal, patients should be counseled about the risk of malignant degeneration in or near the anal transition zone, and can be offered periodic surveillance by endoscopic or anoscopic means every few years, or when symptomatic. ^{274,277,278}

3. Pouchitis is common after IPAA and is managed with antibiotics in most circumstances. Grade of Recommendation: Strong recommendation based on moderate-quality evidence, 1B.

The most frequent long-term complication after IPAA for UC is a nonspecific inflammation of the ileal pouch known as pouchitis. 137,143,279,280 The presence of extraintestinal manifestations of UC before colectomy, especially primary sclerosing cholangitis, has been associated with an increased incidence of pouchitis.^{281,282} It is unclear whether the presence of backwash ileitis or the extent of disease predicts the likelihood of ultimately developing pouchitis, 283-285 although a recent study found the presence of backwash ileitis to be an independent risk factor for chronic pouchitis.²⁸² The origin of this nonspecific inflammation is unclear, but it may be the result of an overgrowth of anaerobic bacteria. 286,287 Presenting symptoms usually include abdominal cramps, fever, pelvic pain, and an increase in stool frequency. Clinical diagnosis may require confirmation by endoscopy and pouch mucosal biopsy.^{288,289} However, it seems that histologic evaluation may be omitted without compromising diagnostic accuracy, and therapy can be successful based on the appearance of the pouch and the appropriate clinical picture.²⁹⁰

Treatment of pouchitis relies primarily on antibiotics, such as metronidazole and ciprofloxacin.^{291–293} Probiotics have been used successfully in pouch patients to provide

prophylaxis against pouchitis and to maintain remission in chronic pouchitis.^{294,295} In a recent Cochrane review, probiotics were found to be more effective than placebo in the treatment of chronic pouchitis as well as the prevention of its onset.²⁹⁶ In antibiotic-refractory cases, budesonide enemas or other medical treatments may be useful.²⁹⁷ Patients who have chronic pouchitis should be assessed for Crohn's disease. Uncommonly, an ileostomy with or without pouch excision is required for severe refractory pouchitis.²⁹²

Ileal inflammation extending proximal to the pouch in the neoterminal ileum has been recently described and termed prepouch ileitis. This nonspecific inflammation may extend up to 50 cm, and biopsies have shown histology similar to pouchitis.²⁹⁸ In a recent study, the incidence of prepouch ileitis was 5.7%.²⁹⁹ All patients had concurrent pouchitis, but less than one-fourth (23.5%) were symptomatic. Early data show no evidence for Crohn's disease in these patients.

APPENDIX A: CONTRIBUTING MEMBERS OF THE ASCRS STANDARDS COMMITTEE

Janice Rafferty, Chair; Scott Steele, Co-Chair; Jose Guillem, Council Representative; W. Donald Buie, Advisor; Andreas Kaiser; George Chang; Dan Feingold; Dan Herzig; John Monson; Scott Strong; Kirsten Wilkins; Marty Weiser; Samantha Hendron; Ian Paquette.

REFERENCES

- Sands BE. Fulminant colitis. J Gastrointest Surg. 2008;12:2157–2159.
- Pal S, Sahni P, Pande GK, Acharya SK, Chattopadhyay TK.
 Outcome following emergency surgery for refractory severe ulcerative colitis in a tertiary care centre in India. BMC Gastroenterol. 2005;5:39.
- 3. Cima RR. Timing and indications for colectomy in chronic ulcerative colitis: Surgical consideration. *Dig Dis.* 2010;28: 501–507.
- 4. Goligher JC, Hoffman DC, de Dombal FT. Surgical treatment of severe attacks of ulcerative colitis, with special reference to the advantages of early operation. *Br Med J.* 1970;4: 703–706.
- 5. Hultén L. Proctocolectomy and ileostomy to pouch surgery for ulcerative colitis. *World J Surg.* 1998;22:335–341.
- 6. Holubar SD, Larson DW, Dozois EJ, Pattana-Arun J, Pemberton JH, Cima RR. Minimally invasive subtotal colectomy and ileal pouch-anal anastomosis for fulminant ulcerative colitis: a reasonable approach? *Dis Colon Rectum.* 2009;52:187–192.
- Holubar SD, Privitera A, Cima RR, Dozois EJ, Pemberton JH, Larson DW. Minimally invasive total proctocolectomy with Brooke ileostomy for ulcerative colitis. *Inflamm Bowel Dis*. 2009;15:1337–1342.
- 8. Cohen JL, Strong SA, Hyman NH, et al; Standards Practice Task Force American Society of Colon and Rectal Surgeons. Practice

- parameters for the surgical treatment of ulcerative colitis. *Dis Colon Rectum.* 2005;48:1997–2009.
- 9. Guyatt G, Gutermen D, Baumann MH, et al. Grading strength of recommendations and quality of evidence in clinical guidelines: report from an American College of Chest Physicians Task Force *Chest.* 2006;129:174–181.
- 10. Truelove SC, Witts LF. Cortisone in ulcerative colitis: final report on a therapeutic trial. *BMJ*. 1955;2:1041–1048.
- 11. Hanauer SB. Inflammatory bowel disease. N Engl J Med. 1996;334:841–848.
- 12. Jones JH, Chapman M. Definition of megacolon in colitis. *Gut.* 1969;10:562–564.
- 13. Katz JA. Medical and surgical management of severe colitis. Semin Gastrointest Dis. 2000;11:18–32.
- 14. Present DH. Toxic megacolon. Med Clin North Am. 1993;77:1129–1148.
- 15. Hyman NH, Cataldo P, Osler T. Urgent subtotal colectomy for severe inflammatory bowel disease. *Dis Colon Rectum*. 2005;48:70–73.
- Greenstein AJ, Barth JA, Sachar DB, Aufses AH Jr. Free colonic perforation without dilatation in ulcerative colitis. *Am J Surg*. 1986;152:272–275.
- Heppell J, Farkouh E, Dubé S, Péloquin A, Morgan S, Bernard D. Toxic megacolon: an analysis of 70 cases. *Dis Colon Rectum*. 1986;29:789–792.
- Greenstein AJ, Sachar DB, Gibas A, et al. Outcome of toxic dilatation in ulcerative and Crohn's colitis. *J Clin Gastroenterol*. 1985;7:137–143.
- Berg DF, Bahadursingh AM, Kaminski DL, Longo WE. Acute surgical emergencies in inflammatory bowel disease. *Am J Surg*. 2002;184:45–51.
- 20. St Peter SD, Abbas MA, Kelly KA. The spectrum of pneumatosis intestinalis. *Arch Surg.* 2003;138:68–75.
- 21. Roy MA. Inflammatory bowel disease. Surg Clin North Am. 1997;77:1419–1436.
- 22. Caprilli R, Latella G, Vernia P, Frieri G. Multiple organ dysfunction in ulcerative colitis. *Am J Gastroenterol.* 2000;95: 1258–1262.
- Bitton A, Buie D, Enns R, et al; Canadian Association of Gastroenterology Severe Ulcerative Colitis Consensus Group. Treatment of hospitalized adult patients with severe ulcerative colitis: Toronto consensus statements. *Am J Gastroenterol*. 2012;107:179–194.
- 24. Oshitani N, Kitano A, Matsumoto T, Kobayashi K. Corticosteroids for the management of ulcerative colitis. *J Gastroenterol.* 1995;30(suppl 8):118–120.
- 25. Sood A, Midha V, Sood N, Awasthi G. A prospective, open-label trial assessing dexamethasone pulse therapy in moderate to severe ulcerative colitis. *J Clin Gastroenterol*. 2002;35:328–331.
- 26. Campbell S, Travis S, Jewell D. Ciclosporin use in acute ulcerative colitis: a long-term experience. *Eur J Gastroenterol Hepatol.* 2005;17:79–84.
- Moskovitz DN, Van Assche G, Maenhout B, et al. Incidence of colectomy during long-term follow-up after cyclosporine-induced remission of severe ulcerative colitis. *Clin Gastroenterol Hepatol*. 2006;4:760–765.
- 28. Bass EM, Del Pino A, Tan A, Pearl RK, Orsay CP, Abcarian H. Does preoperative stoma marking and education by the enterostomal therapist affect outcome? *Dis Colon Rectum.* 1997;40:440–442.

- Person B, Ifargan R, Lachter J, Duek SD, Kluger Y, Assalia A. The impact of preoperative stoma site marking on the incidence of complications, quality of life, and patient's independence. *Dis Colon Rectum.* 2012;55:783–787.
- Järnerot G, Hertervig E, Friis-Liby I, et al. Infliximab as rescue therapy in severe to moderately severe ulcerative colitis: a randomized, placebo-controlled study. *Gastroenterology*. 2005;128:1805–1811.
- 31. Sands BE, Tremaine WJ, Sandborn WJ, et al. Infliximab in the treatment of severe, steroid-refractory ulcerative colitis: a pilot study. *Inflamm Bowel Dis.* 2001;7:83–88.
- 32. Probert CS, Hearing SD, Schreiber S, et al. Infliximab in moderately severe glucocorticoid resistant ulcerative colitis: a randomised controlled trial. *Gut.* 2003;52:998–1002.
- 33. Aratari A, Papi C, Clemente V, et al. Colectomy rate in acute severe ulcerative colitis in the infliximab era. *Dig Liver Dis.* 2008;40:821–826.
- Yamamoto-Furusho JK, Uzcanga LF. Infliximab as a rescue therapy for hospitalized patients with severe ulcerative colitis refractory to systemic corticosteroids. *Dig Surg.* 2008;25:383–386.
- Sandborn WJ, Rutgeerts P, Feagan BG, et al. Colectomy rate comparison after treatment of ulcerative colitis with placebo or infliximab. *Gastroenterology*. 2009;137:1250–1260.
- 36. Regueiro M, Curtis J, Plevy S. Infliximab for hospitalized patients with severe ulcerative colitis. *J Clin Gastroenterol.* 2006;40:476–481.
- 37. Bressler B, Law JK, Al Nahdi Sheraisher N, et al. The use of infliximab for treatment of hospitalized patients with acute severe ulcerative colitis. *Can J Gastroenterol.* 2008;22:937–940.
- 38. Sjöberg M, Magnuson A, Björk J, et al and the Swedish Organization for the Study of Inflammatory Bowel Disease (SOIBD). Infliximab as rescue therapy in hospitalised patients with steroid-refractory acute ulcerative colitis: a long-term follow-up of 211 Swedish patients. *Aliment Pharmacol Ther.* 2013;38:377–387.
- Lichtiger S, Present DH, Kornbluth A, et al. Cyclosporine in severe ulcerative colitis refractory to steroid therapy. N Engl J Med. 1994;330:1841–1845.
- 40. D'Haens G, Lemmens L, Geboes K, et al. Intravenous cyclosporine versus intravenous corticosteroids as single therapy for severe attacks of ulcerative colitis. *Gastroenterology*. 2001;120:1323–1329.
- 41. Shibolet O, Regushevskaya E, Brezis M, Soares-Weiser K. Cyclosporine A for induction of remission in severe ulcerative colitis. *Cochrane Database Syst Rev.* 2005;(1):CD004277.
- 42. Van Assche G, Vermeire S, Rutgeerts P. Management of acute severe ulcerative colitis. *Gut.* 2011;60:130–133.
- 43. Maser EA, Deconda D, Lichtiger S, Ullman T, Present DH, Kornbluth A. Cyclosporine and infliximab as rescue therapy for each other in patients with steroid-refractory ulcerative colitis. *Clin Gastroenterol Hepatol.* 2008;6:1112–1116.
- 44. Leblanc S, Allez M, Seksik P, et al. Successive treatment with cyclosporine and infliximab in severe ulcerative colitis (UC). *Gastroenterology*. 2009; 136(suppl 1):561.
- 45. Mañosa M, López San Román A, Garcia-Planella E, et al. Infliximab rescue therapy after cyclosporin failure in steroid-refractory ulcerative colitis. *Digestion*. 2009;80:30–35.
- 46. Travis SP, Farrant JM, Ricketts C, et al. Predicting outcome in severe ulcerative colitis. *Gut.* 1996;38:905–910.

- 47. Latella G, Vernia P, Viscido A, et al. GI distension in severe ulcerative colitis. *Am J Gastroenterol.* 2002;97:1169–1175.
- 48. Kaplan GG, McCarthy EP, Ayanian JZ, Korzenik J, Hodin R, Sands BE. Impact of hospital volume on postoperative morbidity and mortality following a colectomy for ulcerative colitis. *Gastroenterology*. 2008;134:680–687.
- 49. Roberts SE, Williams JG, Yeates D, Goldacre MJ. Mortality in patients with and without colectomy admitted to hospital for ulcerative colitis and Crohn's disease: record linkage studies. *BMJ*. 2007;335:1033.
- 50. Berger M, Gribetz D, Korelitz BI. Growth retardation in children with ulcerative colitis: the effect of medical and surgical therapy. *Pediatrics*. 1975;55:459–467.
- 51. Griffiths AM. Growth retardation in early-onset inflammatory bowel disease: should we monitor and treat these patients differently? *Dig Dis.* 2009;27:404–411.
- 52. Heuschkel R, Salvestrini C, Beattie RM, Hildebrand H, Walters T, Griffiths A. Guidelines for the management of growth failure in childhood inflammatory bowel disease. *Inflamm Bowel Dis.* 2008;14:839–849.
- Ekbom A, Helmick C, Zack M, Adami HO. Ulcerative colitis and colorectal cancer: a population-based study. N Engl J Med. 1990;323:1228–1233.
- 54. Triantafillidis JK, Nasioulas G, Kosmidis PA. Colorectal cancer and inflammatory bowel disease: epidemiology, risk factors, mechanisms of carcinogenesis and prevention strategies. *Anticancer Res.* 2009;29:2727–2737.
- 55. Askling J, Dickman PW, Karlén P, et al. Family history as a risk factor for colorectal cancer in inflammatory bowel disease. *Gastroenterology*. 2001;120:1356–1362.
- Broomé U, Lindberg G, Löfberg R. Primary sclerosing cholangitis in ulcerative colitis: a risk factor for the development of dysplasia and DNA aneuploidy? *Gastroenterology*. 1992;102:1877–1880.
- 57. Kornfeld D, Ekbom A, Ihre T. Is there an excess risk for colorectal cancer in patients with ulcerative colitis and concomitant primary sclerosing cholangitis? A population based study. *Gut.* 1997;41:522–525.
- Jayaram H, Satsangi J, Chapman RW. Increased colorectal neoplasia in chronic ulcerative colitis complicated by primary sclerosing cholangitis: fact or fiction? *Gut.* 2001;48:430–434.
- Eaden JA, Abrams KR, Mayberry JF. The risk of colorectal cancer in ulcerative colitis: a meta-analysis. Gut. 2001;48:526–535.
- 60. Bernstein CN, Blanchard JF, Kliewer E, Wajda A. Cancer risk in patients with inflammatory bowel disease: a population-based study. *Cancer*. 2001;91:854–862.
- 61. Lakatos L, Mester G, Erdelyi Z, et al. Risk factors for ulcerative colitis-associated colorectal cancer in a Hungarian cohort of patients with ulcerative colitis: results of a population-based study. *Inflamm Bowel Dis.* 2006;12:205–211.
- 62. Collins PD, Mpofu C, Watson AJ, Rhodes JM. Strategies for detecting colon cancer and/or dysplasia in patients with inflammatory bowel disease. *Cochrane Database Syst Rev.* 2006;(2):CD000279.
- 63. Itzkowitz SH, Present DH; Crohn's and Colitis Foundation of America Colon Cancer in IBD Study Group. Consensus conference: colorectal cancer screening and surveillance in inflammatory bowel disease. *Inflamm Bowel Dis.* 2005;11:314–321.
- 64. Riddell RH. Screening strategies in gastrointestinal cancer. *Scand J Gastroenterol Suppl.* 1990;175:177–184.

- 65. Winawer S, Fletcher R, Rex D, et al. Colorectal cancer screening and surveillance: clinical guidelines and rationale—update based on new evidence. *Gastroenterology*. 2003; 124:544–560.
- 66. Kornbluth A, Sachar DB; Practice Parameters Committee of the American College of Gastroenterology. Ulcerative colitis practice guidelines in adults (update): American College of Gastroenterology, Practice Parameters Committee. Am J Gastroenterol. 2004;99:1371–1385.
- 67. Farraye FA, Odze RD, Eaden J, Itzkowitz SH. AGA medical position statement on the diagnosis of colorectal neoplasia in inflammatory bowel disease. *Gastroenterology*. 2010;138:738–745.
- 68. Kornbluth A, Sachar DB; Practice Parameters Committee of the American College of Gastroenterology. Ulcerative colitis practice guidelines in adults: American College of Gastroenterology, Practice Parameters Committee. *Am J Gastroenterol.* 2010;105:501–523.
- 69. Leighton JA, Shen B, Baron TH, et al; Standards of Practice Committee, American Society for Gastrointestinal Endoscopy. ASGE guideline: endoscopy in the diagnosis and treatment of inflammatory bowel disease. *Gastrointest Endosc.* 2006;63:558–565.
- Awais D, Siegel CA, Higgins PD. Modelling dysplasia detection in ulcerative colitis: clinical implications of surveillance intensity. *Gut.* 2009;58:1498–1503.
- 71. Rubin CE, Haggitt RC, Burmer GC, et al. DNA aneuploidy in colonic biopsies predicts future development of dysplasia in ulcerative colitis. *Gastroenterology*. 1992;103:1611–1620.
- Blonski W, Kundu R, Lewis J, Aberra F, Osterman M, Lichtenstein GR. Is dysplasia visible during surveillance colonoscopy in patients with ulcerative colitis? *Scand J Gastroenterol*. 2008;43:698–703.
- Rubin DT, Rothe JA, Hetzel JT, Cohen RD, Hanauer SB. Are dysplasia and colorectal cancer endoscopically visible in patients with ulcerative colitis? *Gastrointest Endosc.* 2007;65:998–1004.
- 74. Marion JF, Waye JD, Present DH, et al; Chromoendoscopy Study Group at Mount Sinai School of Medicine. Chromoendoscopytargeted biopsies are superior to standard colonoscopic surveillance for detecting dysplasia in inflammatory bowel disease patients: a prospective endoscopic trial. *Am J Gastroenterol*. 2008;103:2342–2349.
- 75. Hurlstone DP, Kiesslich R, Thomson M, Atkinson R, Cross SS. Confocal chromoscopic endomicroscopy is superior to chromoscopy alone for the detection and characterisation of intraepithelial neoplasia in chronic ulcerative colitis. *Gut.* 2008;57:196–204.
- Rutter MD, Saunders BP, Schofield G, Forbes A, Price AB, Talbot IC. Pancolonic indigo carmine dye spraying for the detection of dysplasia in ulcerative colitis. *Gut.* 2004;53:256–260.
- 77. Hurlstone DP, Sanders DS, Lobo AJ, McAlindon ME, Cross SS. Indigo carmine-assisted high-magnification chromoscopic colonoscopy for the detection and characterisation of intraepithelial neoplasia in ulcerative colitis: a prospective evaluation. *Endoscopy.* 2005;37:1186–1192.
- 78. Kiesslich R, Goetz M, Lammersdorf K, et al. Chromoscopyguided endomicroscopy increases the diagnostic yield of intraepithelial neoplasia in ulcerative colitis. *Gastroenterology*. 2007;132:874–882.
- 79. Itzkowitz SH, Yio X. Inflammation and cancer IV. Colorectal cancer in inflammatory bowel disease: the role of inflammation. *Am J Physiol Liver Physiol*. 2004;287:G7–G17.

- 80. Breynaert C, Vermeire S, Rutgeerts P, Van Assche G. Dysplasia and colorectal cancer in inflammatory bowel disease: a result of inflammation or an intrinsic risk? *Acta Gastroenterol Belg.* 2008:71:367–372.
- 81. Shapiro BD, Lashner BA. Cancer biology in ulcerative colitis and potential use in endoscopic surveillance. *Gastrointest Endosc Clin N Am.* 1997;7:453–468.
- 82. Bernstein CN, Shanahan F, Weinstein WM. Are we telling patients the truth about surveillance colonoscopy in ulcerative colitis? *Lancet*. 1994;343:71–74.
- 83. Blackstone MO, Riddell RH, Rogers BH, Levin B. Dysplasia-associated lesion or mass (DALM) detected by colonoscopy in long-standing ulcerative colitis: an indication for colectomy. *Gastroenterology*. 1981;80:366–374.
- 84. Riddell RH, Goldman H, Ransohoff DF, et al. Dysplasia in inflammatory bowel disease: standardized classification with provisional clinical applications. *Hum Pathol.* 1983;14:931–968.
- 85. Torres C, Antonioli D, Odze RD. Polypoid dysplasia and adenomas in inflammatory bowel disease: a clinical, pathologic, and follow-up study of 89 polyps from 59 patients. *Am J Surg Pathol.* 1998;22:275–284.
- 86. Blonski W, Kundu R, Furth EF, Lewis J, Aberra F, Lichtenstein GR. High-grade dysplastic adenoma-like mass lesions are not an indication for colectomy in patients with ulcerative colitis. *Scand J Gastroenterol.* 2008;43:817–820.
- 87. Odze RD, Farraye FA, Hecht JL, Hornick JL. Long-term followup after polypectomy for adenoma-like dysplastic lesions in ulcerative colitis. *Clin Gastroenterol Hepatol.* 2004;2:534–541.
- 88. Medlicott SA, Jewell LD, Price L, Fedorak RN, Sherbaniuk RW, Urbanski SJ. Conservative management of small adenomata in ulcerative colitis. *Am J Gastroenterol.* 1997;92:2094–2098.
- 89. Rubin PH, Friedman S, Harpaz N, et al. Colonoscopic polypectomy in chronic colitis: conservative management after endoscopic resection of dysplastic polyps. *Gastroenterology*. 1999;117:1295–1300.
- 90. Friedman S, Odze RD, Farraye FA. Management of neoplastic polyps in inflammatory bowel disease. *Inflamm Bowel Dis.* 2003;9:260–266.
- 91. Itzkowitz SH, Harpaz N. Diagnosis and management of dysplasia in patients with inflammatory bowel diseases. *Gastroenterology*. 2004;126:1634–1648.
- 92. Befrits R, Ljung T, Jaramillo E, Rubio C. Low-grade dysplasia in extensive, long-standing inflammatory bowel disease: a follow-up study. *Dis Colon Rectum*. 2002;45:615–620.
- 93. Lim CH, Dixon MF, Vail A, Forman D, Lynch DA, Axon AT. Ten year follow up of ulcerative colitis patients with and without low grade dysplasia. *Gut.* 2003;52:1127–1132.
- Connell WR, Lennard-Jones JE, Williams CB, Talbot IC, Price AB, Wilkinson KH. Factors affecting the outcome of endoscopic surveillance for cancer in ulcerative colitis. *Gastroenterology*. 1994;107:934–944.
- 95. Ullman T, Croog V, Harpaz N, Sachar D, Itzkowitz S. Progression of flat low-grade dysplasia to advanced neoplasia in patients with ulcerative colitis. *Gastroenterology*. 2003;125:1311–1319.
- 96. Thomas T, Abrams KA, Robinson RJ, Mayberry JF. Metaanalysis: cancer risk of low-grade dysplasia in chronic ulcerative colitis. *Aliment Pharmacol Ther*. 2007;25:657–668.
- 97. Eaden J, Abrams K, Ekbom A, Jackson E, Mayberry J. Colorectal cancer prevention in ulcerative colitis: a case-control study. *Aliment Pharmacol Ther.* 2000;14:145–153.

- 98. Odze RD, Goldblum J, Noffsinger A, Alsaigh N, Rybicki LA, Fogt F. Interobserver variability in the diagnosis of ulcerative colitis-associated dysplasia by telepathology. *Mod Pathol.* 2002;15:379–386.
- 99. Gorfine SR, Bauer JJ, Harris MT, Kreel I. Dysplasia complicating chronic ulcerative colitis: is immediate colectomy warranted? *Dis Colon Rectum.* 2000;43:1575–1581.
- 100. Ullman TA, Loftus EV Jr, Kakar S, Burgart LJ, Sandborn WJ, Tremaine WJ. The fate of low grade dysplasia in ulcerative colitis. *Am J Gastroenterol.* 2002;97:922–927.
- 101. Velayos FS, Terdiman JP, Walsh JM. Effect of 5-aminosalicylate use on colorectal cancer and dysplasia risk: a systematic review and metaanalysis of observational studies. *Am J Gastroenterol.* 2005;100:1345–1353.
- 102. Rubin DT, LoSavio A, Yadron N, Huo D, Hanauer SB. Aminosalicylate therapy in the prevention of dysplasia and colorectal cancer in ulcerative colitis. Clin Gastroenterol Hepatol. 2006;4:1346–1350.
- 103. van Staa TP, Card T, Logan RF, Leufkens HG. 5-Aminosalicylate use and colorectal cancer risk in inflammatory bowel disease: a large epidemiological study. *Gut.* 2005;54:1573–1578.
- 104. Terdiman JP, Steinbuch M, Blumentals WA, Ullman TA, Rubin DT. 5-Aminosalicylic acid therapy and the risk of colorectal cancer among patients with inflammatory bowel disease. *Inflamm Bowel Dis.* 2007;13:367–371.
- 105. Bernstein CN, Nugent Z, Blanchard JF. 5-Aminosalicylate is not chemoprophylactic for colorectal cancer in IBD: a population based study. *Am J Gastroenterol*. 2011;106:731–736.
- 106. Lashner BA, Turner BC, Bostwick DG, Frank PH, Hanauer SB. Dysplasia and cancer complicating strictures in ulcerative colitis. *Dig Dis Sci.* 1990;35:349–352.
- 107. Gumaste V, Sachar DB, Greenstein AJ. Benign and malignant strictures in ulcerative colitis. *Gut.* 1992;33:938–941.
- 108. Reiser JR, Waye JD, Janowitz HD, Harpaz N. Adenocarcinoma in strictures of ulcerative colitis without antecedent dysplasia by colonoscopy. *Am J Gastroenterol*. 1994;89:119–122.
- 109. Alves A, Panis Y, Bouhnik Y, Maylin V, Lavergne-Slove A, Valleur P. Subtotal colectomy for severe acute colitis: a 20-year experience of a tertiary care center with an aggressive and early surgical policy. *J Am Coll Surg.* 2003;197:379–385.
- 110. Penna C, Daude F, Parc R, et al. Previous subtotal colectomy with ileostomy and sigmoidostomy improves the morbidity and early functional results after ileal pouch-anal anastomosis in ulcerative colitis. *Dis Colon Rectum.* 1993;36:343–348.
- 111. Ouaïssi M, Alves A, Bouhnik Y, Valleur P, Panis Y. Three-step ileal pouch-anal anastomosis under total laparoscopic approach for acute or severe colitis complicating inflammatory bowel disease. *J Am Coll Surg.* 2006;202:637–642.
- 112. Marceau C, Alves A, Ouaissi M, Bouhnik Y, Valleur P, Panis Y. Laparoscopic subtotal colectomy for acute or severe colitis complicating inflammatory bowel disease: a case-matched study in 88 patients. *Surgery*. 2007;141:640–644.
- 113. Dunker MS, Bemelman WA, Slors JF, van Hogezand RA, Ringers J, Gouma DJ. Laparoscopic-assisted vs open colectomy for severe acute colitis in patients with inflammatory bowel disease (IBD): a retrospective study in 42 patients. *Surg Endosc.* 2000;14:911–914.
- 114. Carter FM, McLeod RS, Cohen Z. Subtotal colectomy for ulcerative colitis: complications related to the rectal remnant. *Dis Colon Rectum.* 1991;34:1005–1009.

- 115. Karch LA, Bauer JJ, Gorfine SR, Gelernt IM. Subtotal colectomy with Hartmann's pouch for inflammatory bowel disease. *Dis Colon Rectum.* 1995;38:635–639.
- 116. Ziv Y, Fazio VW, Church JM, Milsom JW, Schroeder TK. Safety of urgent restorative proctocolectomy with ileal pouchanal anastomosis for fulminant colitis. *Dis Colon Rectum*. 1995;38:345–349.
- 117. Harms BA, Myers GA, Rosenfeld DJ, Starling JR. Management of fulminant ulcerative colitis by primary restorative proctocolectomy. *Dis Colon Rectum.* 1994;37:971–978.
- 118. Jimmo B, Hyman NH. Is ileal pouch-anal anastomosis really the procedure of choice for patients with ulcerative colitis? *Dis Colon Rectum.* 1998;41:41–45.
- 119. Fazio VW, Tekkis PP, Remzi F, et al. Quantification of risk for pouch failure after ileal pouch anal anastomosis surgery. *Ann Surg.* 2003;238:605–614.
- 120. Metcalf AM. Elective and emergent operative management of ulcerative colitis. *Surg Clin North Am.* 2007;87:633–641.
- 121. Causey MW, Stoddard D, Johnson EK, et al. Laparoscopy impacts outcomes favorably following colectomy for ulcerative colitis: a critical analysis of the ACS-NSQIP database. *Surg Endosc.* 2013;27:603–609.
- 122. Carlstedt A, Fasth S, Hultén L, Nordgren S, Palselius I. Longterm ileostomy complications in patients with ulcerative colitis and Crohn's disease. *Int J Colorectal Dis.* 1987;2:22–25.
- 123. Camilleri-Brennan J, Munro A, Steele RJ. Does an ileoanal pouch offer a better quality of life than a permanent ileostomy for patients with ulcerative colitis? *J Gastrointest Surg.* 2003;7:814–819.
- 124. Wikland M, Jansson I, Asztély M, et al. Gynaecological problems related to anatomical changes after conventional proctocolectomy and ileostomy. *Int J Colorectal Dis.* 1990;5:49–52.
- 125. Metcalf AM, Dozois RR, Kelly KA. Sexual function in women after proctocolectomy. *Ann Surg.* 1986;204:624–627.
- Yamaguchi N, Isomoto H, Shikuwa S, et al. Proximal extension of backwash ileitis in ulcerative-colitis-associated colon cancer. *Med Sci Monit.* 2010;16:CS87–CS91.
- 127. Annam V, Panduranga C, Kodandaswamy C, Suresh DR. Primary mucinous adenocarcinoma in an ileostomy with adjacent skin invasion: a late complication of surgery for ulcerative colitis. *J Gastrointest Cancer*. 2008;39:138–140.
- 128. Metzger PP, Slappy AL, Chua HK, Menke DM. Adenocarcinoma developing at an ileostomy: report of a case and review of the literature. *Dis Colon Rectum*. 2008;51:604–609.
- 129. Pedersen ME, Rahr HB, Fenger C, Qvist N. Adenocarcinoma arising from the rectal stump eleven years after excision of an ileal J-pouch in a patient with ulcerative colitis: report of a case. *Dis Colon Rectum.* 2008;51:1146–1148.
- 130. Hahnloser D, Pemberton JH, Wolff BG, Larson DR, Crownhart BS, Dozois RR. The effect of aging on function and quality of life in ileal pouch patients: a single cohort experience of 409 patients with chronic ulcerative colitis. *Ann Surg.* 2004; 240:615–621.
- 131. McIntyre PB, Pemberton JH, Wolff BG, Beart RW, Dozois RR. Comparing functional results one year and ten years after ileal pouch-anal anastomosis for chronic ulcerative colitis. *Dis Colon Rectum.* 1994;37:303–307.
- 132. Leowardi C, Hinz U, Tariverdian M, et al. Long-term outcome 10 years or more after restorative proctocolectomy and

- ileal pouch-anal anastomosis in patients with ulcerative colitis. *Langenbecks Arch Surg.* 2010;395:49–56.
- 133. Boller AM, Larson DW. Laparoscopic restorative proctocolectomy for ulcerative colitis. *J Gastrointest Surg.* 2007;11:3–7.
- 134. Larson DW, Dozois EJ, Piotrowicz K, Cima RR, Wolff BG, Young-Fadok TM. Laparoscopic-assisted vs. open ileal pouchanal anastomosis: functional outcome in a case-matched series. *Dis Colon Rectum.* 2005;48:1845–1850.
- 135. Tsuruta M, Hasegawa H, Ishii Y, et al. Hand-assisted versus conventional laparoscopic restorative proctocolectomy for ulcerative colitis. Surg Laparosc Endosc Percutan Tech. 2009;19:52–56.
- 136. Fazio VW, Ziv Y, Church JM, et al. Ileal pouch-anal anastomoses complications and function in 1005 patients. *Ann Surg.* 1995;222:120–127.
- 137. Meagher AP, Farouk R, Dozois RR, Kelly KA, Pemberton JH. J ileal pouch-anal anastomosis for chronic ulcerative colitis: complications and long-term outcome in 1310 patients. *Br J Surg.* 1998;85:800–818.
- 138. Selvaggi F, Sciaudone G, Limongelli P, et al. The effect of pelvic septic complications on function and quality of life after ileal pouch-anal anastomosis: a single center experience. *Am Surg.* 2010;76:428–435.
- 139. Martin A, Dinca M, Leone L, et al. Quality of life after proctocolectomy and ileo-anal anastomosis for severe ulcerative colitis. *Am J Gastroenterol.* 1998;93:166–169.
- 140. Tiainen J, Matikainen M. Health-related quality of life after ileal J-pouch-anal anastomosis for ulcerative colitis: long-term results. *Scand J Gastroenterol*. 1999;34:601–605.
- 141. Carmon E, Keidar A, Ravid A, Goldman G, Rabau M. The correlation between quality of life and functional outcome in ulcerative colitis patients after proctocolectomy ileal pouch anal anastomosis. *Colorectal Dis.* 2003;5:228–232.
- 142. Scarpa M, Angriman I, Ruffolo C, et al. Health-related quality of life after restorative proctocolectomy for ulcerative colitis: long-term results. *World J Surg.* 2004;28:124–129.
- 143. Ferrante M, Declerck S, De Hertogh G, et al. Outcome after proctocolectomy with ileal pouch-anal anastomosis for ulcerative colitis. *Inflamm Bowel Dis.* 2008;14:20–28.
- 144. Cornish JA, Tan E, Teare J, et al. The effect of restorative proctocolectomy on sexual function, urinary function, fertility, pregnancy and delivery: a systematic review. *Dis Colon Rectum*. 2007;50:1128–1138.
- 145. Ørding Olsen K, Juul S, Berndtsson I, Oresland T, Laurberg S. Ulcerative colitis: female fecundity before diagnosis, during disease, and after surgery compared with a population sample. *Gastroenterology*. 2002;122:15–19.
- 146. Johnson P, Richard C, Ravid A, et al. Female infertility after ileal pouch-anal anastomosis for ulcerative colitis. *Dis Colon Rectum*. 2004;47:1119–1126.
- 147. Gorgun E, Remzi FH, Goldberg JM, et al. Fertility is reduced after restorative proctocolectomy with ileal pouch anal anastomosis: a study of 300 patients. *Surgery*. 2004;136:795–803.
- 148. Araki Y, Ishibashi N, Ogata Y, Shirouzu K, Isomoto H. The usefulness of restorative laparoscopic-assisted total colectomy for ulcerative colitis. *Kurume Med J.* 2001;48:99–103.
- 149. Ky AJ, Sonoda T, Milsom JW. One-stage laparoscopic restorative proctocolectomy: an alternative to the conventional approach? *Dis Colon Rectum*. 2002;45:207–211.
- 150. Hasegawa H, Watanabe M, Baba H, Nishibori H, Kitajima M. Laparoscopic restorative proctocolectomy for patients

- with ulcerative colitis. *J Laparoendosc Adv Surg Tech A*. 2002;12:403–406.
- 151. Pace DE, Seshadri PA, Chiasson PM, Poulin EC, Schlachta CM, Mamazza J. Early experience with laparoscopic ileal pouch-anal anastomosis for ulcerative colitis. *Surg Laparosc Endosc Percutan Tech.* 2002;12:337–341.
- 152. Maartense S, Dunker MS, Slors JF, et al. Hand-assisted laparoscopic versus open restorative proctocolectomy with ileal pouch-anal anastomosis: a randomized trial. *Ann Surg* 2004; 240:984–991.
- 153. Kienle P, Z'graggen K, Schmidt J, Benner A, Weitz J, Büchler MW. Laparoscopic restorative proctocolectomy. *Br J Surg.* 2005;92:88–93.
- 154. Alexander F. Complications of ileal pouch anal anastomosis. *Semin Pediatr Surg.* 2007;16:200–204.
- 155. Maruthachalam K, Kumar R, Hainsworth P. Parking the pouch: pouch salvage after anastomotic leak following restorative proctocolectomy: report of a case. *Dis Colon Rectum*. 2008;51:1724–1726.
- 156. Mathis KL, Dozois EJ, Larson DW, Cima RR, Wolff BG, Pemberton JH. Outcomes in patients with ulcerative colitis undergoing partial or complete reconstructive surgery for failing ileal pouch-anal anastomosis. *Ann Surg.* 2009;249:409–413.
- 157. Ahmed Ali U, Keus F, Heikens JT, et al. Open versus laparoscopic (assisted) ileo pouch anal anastomosis for ulcerative colitis and familial adenomatous polyposis. *Cochrane Database Syst Rev.* 2009;(1):CD006267.
- 158. Fajardo AD, Dharmarajan S, George V, et al. Laparoscopic versus open 2-stage ileal pouch: laparoscopic approach allows for faster restoration of intestinal continuity. *J Am Coll Surg.* 2010;211:377–383.
- 159. Hull TL, Joyce MR, Geisler DP, Coffey JC. Adhesions after laparoscopic and open ileal pouch-anal anastomosis surgery for ulcerative colitis. *Br J Surg.* 2012;99:270–275.
- 160. Indar AA, Efron JE, Young-Fadok TM. Laparoscopic ileal pouch-anal anastomosis reduces abdominal and pelvic adhesions. *Surg Endosc.* 2009;23:174–177.
- 161. Lake JP, Firoozmand E, Kang JC, et al. Effect of high-dose steroids on anastomotic complications after proctocolectomy with ileal pouch-anal anastomosis. *J Gastrointest Surg.* 2004;8:547–551.
- 162. Colombel JF, Loftus EV Jr, Tremaine WJ, et al. The safety profile of infliximab in patients with Crohn's disease: the Mayo clinic experience in 500 patients. *Gastroenterology*. 2004;126:19–31.
- 163. Selvasekar CR, Cima RR, Larson DW, et al. Effect of infliximab on short-term complications in patients undergoing operation for chronic ulcerative colitis. *J Am Coll Surg.* 2007;204:956–963.
- 164. Mor IJ, Vogel JD, da Luz Moreira A, Shen B, Hammel J, Remzi FH. Infliximab in ulcerative colitis is associated with an increased risk of postoperative complications after restorative proctocolectomy. *Dis Colon Rectum.* 2008;51:1202–1210.
- 165. Schluender SJ, Ippoliti A, Dubinsky M, et al. Does infliximab influence surgical morbidity of ileal pouch-anal anastomosis in patients with ulcerative colitis? *Dis Colon Rectum*. 2007;50:1747–1753.
- 166. Coquet-Reinier B, Berdah SV, Grimaud JC, et al. Preoperative infliximab treatment and postoperative complications after laparoscopic restorative proctocolectomy with ileal pouch-anal anastomosis: a case-matched study. *Surg Endosc.* 2010;24:1866–1871.

- 167. Bordeianou L, Kunitake H, Shellito P, Hodin R. Preoperative infliximab treatment in patients with ulcerative and indeterminate colitis does not increase rate of conversion to emergent and multistep abdominal surgery. *Int J Colorectal Dis.* 2010;25:401–404.
- 168. Kunitake H, Hodin R, Shellito PC, Sands BE, Korzenik J, Bordeianou L. Perioperative treatment with infliximab in patients with Crohn's disease and ulcerative colitis is not associated with an increased rate of postoperative complications. *J Gastrointest Surg.* 2008;12:1730–1736.
- 169. Hudson M, Flett G, Sinclair TS, Brunt PW, Templeton A, Mowat NA. Fertility and pregnancy in inflammatory bowel disease. Int J Gynaecol Obstet. 1997;58:229–237.
- 170. Olsen KO, Joelsson M, Laurberg S, Oresland T. Fertility after ileal pouch-anal anastomosis in women with ulcerative colitis. *Br J Surg.* 1999;86:493–495.
- 171. Lepistö A, Sarna S, Tiitinen A, Järvinen HJ. Female fertility and childbirth after ileal pouch-anal anastomosis for ulcerative colitis. *Br J Surg*, 2007;94:478–482.
- 172. Waljee A, Waljee J, Morris AM, Higgins PD. Threefold increased risk of infertility: a meta-analysis of infertility after ileal pouch anal anastomosis in ulcerative colitis. *Gut.* 2006; 55:1575–1580.
- 173. Mortier PE, Gambiez L, Karoui M, et al. Colectomy with ileorectal anastomosis preserves female fertility in ulcerative colitis. *Gastroenterol Clin Biol*. 2006;30:594–597.
- 174. Nieuwenhuis MH, Douma KF, Bleiker EM, Bemelman WA, Aaronson NK, Vasen HF. Female fertility after colorectal surgery for familial adenomatous polyposis: a nationwide cross-sectional study. *Ann Surg.* 2010;252:341–344.
- 175. Beyer-Berjot L, Maggiori L, Birnbaum D, Lefevre JH, Berdah S, Panis Y. A total laparoscopic approach reduces the infertility rate after ileal pouch-anal anastomosis: a 2-center study. *Ann Surg.* 2013;258:275–282.
- 176. Bartels SA, D'Hoore A, Cuesta MA, Bensdorp AJ, Lucas C, Bemelman WA. Significantly increased pregnancy rates after laparoscopic restorative proctocolectomy: a cross-sectional study. *Ann Surg.* 2012;256:1045–1048.
- 177. McLeod RS. Ileal pouch anal anastomosis: pregnancy–before, during and after. *J Gastrointest Surg.* 2008;12:2150–2152.
- 178. Seligman NS, Sbar W, Berghella V. Pouch function and gastrointestinal complications during pregnancy after ileal pouch-anal anastomosis. *J Matern Fetal Neonatal Med.* 2001;24:525–530.
- 179. Hahnloser D, Pemberton JH, Wolff BG, et al. Pregnancy and delivery before and after ileal pouch-anal anastomosis for inflammatory bowel disease: immediate and long-term consequences and outcomes. *Dis Colon Rectum*. 2004;47:1127–1135.
- 180. Remzi FH, Gorgun E, Bast J, et al. Vaginal delivery after ileal pouch-anal anastomosis: a word of caution. *Dis Colon Rectum*. 2005;48:1691–1699.
- 181. Gearhart SL, Hull TL, Schroeder T, Church J, Floruta C. Sphincter defects are not associated with long-term incontinence following ileal pouch-anal anastomosis. *Dis Colon Rectum.* 2005;48:1410–1415.
- 182. Kitayama T, Funayama Y, Fukushima K, et al. Anal function during pregnancy and postpartum after ileal pouch anal anastomosis for ulcerative colitis. *Surg Today*. 2005;35:211–215.
- 183. Ravid A, Richard CS, Spencer LM, et al. Pregnancy, delivery, and pouch function after ileal pouch-anal anastomosis for ulcerative colitis. *Dis Colon Rectum*. 2002;45:1283–1288.

- 184. Damgaard B, Wettergren A, Kirkegaard P. Social and sexual function following ileal pouch-anal anastomosis. *Dis Colon Rectum.* 1995;38:286–289.
- 185. Tiainen J, Matikainen M, Hiltunen KM. Ileal J-pouch–anal anastomosis, sexual dysfunction, and fertility. *Scand J Gastroenterol*. 1999;34:185–188.
- 186. Michelassi F, Stella M, Block GE. Prospective assessment of functional results after ileal J pouch-anal restorative proctocolectomy. *Arch Surg.* 1993;128:889–894.
- 187. Berndtsson I, Oresland T, Hultén L. Sexuality in patients with ulcerative colitis before and after restorative proctocolectomy: a prospective study. *Scand J Gastroenterol*. 2004;39:374–379.
- 188. Farouk R, Pemberton JH, Wolff BG, Dozois RR, Browning S, Larson D. Functional outcomes after ileal pouch-anal anastomosis for chronic ulcerative colitis. *Ann Surg*. 2000;231:919–926.
- 189. Gorgun E, Remzi FH, Montague DK, et al. Male sexual function improves after ileal pouch anal anastomosis. *Colorectal Dis.* 2005;7:545–550.
- 190. Larson DW, Davies MM, Dozois EJ, et al. Sexual function, body image, and quality of life after laparoscopic and open ileal pouch-anal anastomosis. *Dis Colon Rectum.* 2008;51:392–396.
- 191. van Balkom KA, Beld MP, Visschers RG, van Gemert WG, Breukink SO. Long-term results after restorative proctocolectomy with ileal pouch-anal anastomosis at a young age. *Dis Colon Rectum.* 2012;55:939–947.
- 192. Zmora O, Spector D, Dotan I, Klausner JM, Rabau M, Tulchinsky H. Is stapled ileal pouch anal anastomosis a safe option in ulcerative colitis patients with dysplasia or cancer? *Int J Colorectal Dis.* 2009;24:1181–1186.
- 193. Taylor BA, Wolff BG, Dozois RR, Kelly KA, Pemberton JH, Beart RW Jr. Ileal pouch-anal anastomosis for chronic ulcerative colitis and familial polyposis coli complicated by adenocarcinoma. *Dis Colon Rectum.* 1988;31:358–362.
- 194. Radice E, Nelson H, Devine RM, et al. Ileal pouch-anal anastomosis in patients with colorectal cancer: long-term functional and oncologic outcomes. *Dis Colon Rectum.* 1998;41:11–17.
- 195. Ziv Y, Fazio VW, Strong SA, Oakley JR, Milsom JW, Lavery IC. Ulcerative colitis and coexisting colorectal cancer: recurrence rate after restorative proctocolectomy. *Ann Surg Oncol*. 1994;1:512–515.
- Gorfine SR, Harris MT, Bub DS, Bauer JJ. Restorative proctocolectomy for ulcerative colitis complicated by colorectal cancer. *Dis Colon Rectum*. 2004;47:1377–1385.
- 197. Stelzner M, Fonkalsrud EW. The endorectal ileal pullthrough procedure in patients with ulcerative colitis and familial polyposis with carcinoma. Surg Gynecol Obstet. 1989;169:187–194.
- 198. Wiltz O, Hashmi HF, Schoetz DJ Jr, et al. Carcinoma and the ileal pouch-anal anastomosis. *Dis Colon Rectum*. 1991;34:805–809.
- 199. Ho KS, Chang CC, Baig MK, et al. Ileal pouch anal anastomosis for ulcerative colitis is feasible for septuagenarians. *Colorectal Dis.* 2006;8:235–238.
- 200. Tan HT, Connolly AB, Morton D, Keighley MR. Results of restorative proctocolectomy in the elderly. *Int J Colorectal Dis.* 1997;12:319–322.
- 201. Takao Y, Gilliland R, Nogueras JJ, Weiss EG, Wexner SD. Is age relevant to functional outcome after restorative proctocolectomy for ulcerative colitis?: prospective assessment of 122 cases. *Ann Surg.* 1998;227:187–194.
- 202. Delaney CP, Fazio VW, Remzi FH, et al. Prospective, agerelated analysis of surgical results, functional outcome, and

- quality of life after ileal pouch-anal anastomosis. *Ann Surg.* 2003;238:221–228.
- 203. Longo WE, Virgo KS, Bahadursingh AN, Johnson FE. Patterns of disease and surgical treatment among United States veterans more than 50 years of age with ulcerative colitis. *Am J Surg.* 2003;186:514–518.
- 204. Michelassi F, Lee J, Rubin M, et al. Long-term functional results after ileal pouch anal restorative proctocolectomy for ulcerative colitis: a prospective observational study. *Ann Surg.* 2003;238:433–441.
- 205. Silvestri MT, Hurst RD, Rubin MA, Michelassi F, Fichera A. Chronic inflammatory changes in the anal transition zone after stapled ileal pouch-anal anastomosis: is mucosectomy a superior alternative? *Surgery*. 2008;144:533–537.
- 206. Gosselink MP, West RL, Kuipers EJ, Hansen BE, Schouten WR. Integrity of the anal sphincters after pouch-anal anastomosis: evaluation with three-dimensional endoanal ultrasonography. *Dis Colon Rectum.* 2005;48:1728–1735.
- 207. Seow-Choen A, Tsunoda A, Nicholls RJ. Prospective randomized trial comparing anal function after handsewn ileoanal anastomosis versus stapled ileoanal anastomosis without mucosectomy in restorative proctocolectomy. *Br J Surg.* 1991; 78:430–434.
- 208. Luukkonen P, Jarvinen H. Stapled versus hand sutured ileoanal anastomosis in restorative proctocolectomy: a prospective randomized trial. *Arch Surg.* 1993;128:437–440.
- 209. Reilly WT, Pemberton JH, Wolff BG, et al. Randomized prospective trial comparing ileal pouch-anal anastomosis performed by excising the anal mucosa to ileal pouch-anal anastomosis. *Ann Surg.* 1997; 225:666–676.
- 210. Bednarz W, Olewinski R, Wojczys R, Sutkowski K, Domoslawski P, Balcerzak W. Ileal-pouch-anal anastomosis after restorative proctocolectomy in patients with ulcerative colitis or familial adenomatous polyposis. *Hepatogastroenterology*. 2005;52:1101–1105.
- 211. Hasegawa S, Nomura A, Kawamura J, et al. Laparoscopic restorative total proctocolectomy with mucosal resection. *Dis Colon Rectum.* 2007;50:1152–1156.
- 212. Utsunomiya J, Iwama T, Imajo M, et al. Total colectomy, mucosal proctectomy, and ileoanal anastomosis. *Dis Colon Rectum*. 1980;23:459–466.
- 213. Fonkalsrud EW. Total colectomy and endorectal ileal pullthrough with internal ileal reservoir for ulcerative colitis. Surg Gynecol Obstet. 1980;150:1–8.
- 214. Nicholls RJ, Lubowski DZ. Restorative proctocolectomy: the four loop (W) reservoir. *Br J Surg.* 1987;74:564–566.
- 215. Johnston D, Williamson ME, Lewis WG, Miller AS, Sagar PM, Holdsworth PJ. Prospective controlled trial of duplicated (J) versus quadruplicated (W) pelvic ileal reservoirs in restorative proctocolectomy for ulcerative colitis. *Gut.* 1996;39:242–247.
- 216. Keighley MR, Yoshioka K, Kmiot W. Prospective randomized trial to compare the stapled double lumen pouch and the sutured quadruple pouch for restorative proctocolectomy. *Br J Surg.* 1988;75:1008–1011.
- 217. Smith L, Friend WG, Medwell SJ. The superior mesenteric artery: the critical factor in the pouch pull-through procedure. *Dis Colon Rectum.* 1984;27:741–744.
- 218. Cohen Z, McLeod RS, Stephen W, Stern HS, O'Connor B, Reznick R. Continuing evolution of the pelvic pouch procedure. *Ann Surg.* 1992;216:506–511.

- 219. Galandiuk S, Wolff BG, Dozois RR, Beart RW Jr. Ileal pouchanal anastomosis without ileostomy. *Dis Colon Rectum*. 1991;34:870–873.
- 220. Gorfine SR, Gelernt IM, Bauer JJ, Harris MT, Kreel I. Restorative proctocolectomy without diverting ileostomy. *Dis Colon Rectum.* 1995;38:188–194.
- 221. Joyce MR, Kiran RP, Remzi FH, Church J, Fazio VW. In a select group of patients meeting strict clinical criteria and undergoing ileal pouch-anal anastomosis, the omission of a diverting ileostomy offers cost savings to the hospital. *Dis Colon Rectum*. 2010;53:905–910.
- 222. Sugerman HJ, Sugerman EL, Meador JG, Newsome HH Jr, Kellum JM Jr, DeMaria EJ. Ileal pouch anal anastomosis without ileal diversion. *Ann Surg.* 2000;232:530–541.
- 223. Lovegrove RE, Symeonides P, Tekkis PP, Goodfellow PB, Shorthouse AJ. A selective approach to restorative proctocolectomy without ileostomy: a single centre experience. *Colorectal Dis.* 2008;10:916–924.
- 224. Remzi FH, Fazio VW, Gorgun E, et al. The outcome after restorative proctocolectomy with or without defunctioning ileostomy. *Dis Colon Rectum*. 2006;49:470–477.
- 225. Tjandra JJ, Fazio VW, Milsom JW, Lavery IC, Oakley JR, Fabre JM. Omission of temporary diversion in restorative procto-colectomy—is it safe? *Dis Colon Rectum*. 1993;36:1007–1014.
- 226. Williamson ME, Lewis WG, Sagar PM, Holdsworth PJ, Johnston D. One-stage restorative proctocolectomy without temporary ileostomy for ulcerative colitis: a note of caution. *Dis Colon Rectum.* 1997;40:1019–1022.
- 227. Davies M, Hawley PR. Ten years experience of one-stage restorative proctocolectomy for ulcerative colitis. *Int J Colorectal Dis.* 2007;22:1255–1260.
- 228. Ikeuchi H, Nakano H, Uchino M, et al. Safety of one-stage restorative proctocolectomy for ulcerative colitis. *Dis Colon Rectum*. 2005;48:1550–1555.
- 229. Gullberg K, Liljeqvist L. Stapled ileoanal pouches without loop ileostomy: a prospective study in 86 patients. *Int J Colorectal Dis.* 2001;16:221–227.
- 230. Farouk R, Dozois RR, Pemberton JH, Larson D. Incidence and subsequent impact of pelvic abscess after ileal pouch-anal anastomosis for chronic ulcerative colitis. *Dis Colon Rectum*. 1998;41:1239–1243.
- 231. Gunnarsson U, Karlbom U, Docker M, Raab Y, Påhlman L. Proctocolectomy and pelvic pouch—is a diverting stoma dangerous for the patient? *Colorectal Dis.* 2004;6:23–27.
- 232. Wong NY, Eu KW. A defunctioning ileostomy does not prevent clinical anastomotic leak after a low anterior resection: a prospective, comparative study. *Dis Colon Rectum*. 2005;48:2076–2079.
- 233. MacLean AR, Cohen Z, MacRae HM, et al. Risk of small bowel obstruction after the ileal pouch-anal anastomosis. *Ann Surg.* 2002;235:200–206.
- 234. Mowschenson PM, Critchlow JF, Peppercorn MA. Ileoanal pouch operation: long-term outcome with or without diverting ileostomy. *Arch Surg.* 2000;135:463–465.
- 235. Lian L, Fazio VW, Remzi FH, Shen B, Dietz D, Kiran RP. Outcomes for patients undergoing continent ileostomy after a failed ileal pouch-anal anastomosis. *Dis Colon Rectum.* 2009;52:1409–1414.
- 236. Hultén L, Fasth S, Hallgren T, Oresland T. The failing pelvic pouch conversion to continent ileostomy. *Int J Colorectal Dis.* 1992;7:119–121.

- 237. Lepisto AH, Jarvinen HJ. Durability of the Kock continent ileostomy. *Dis Colon Rectum*. 2003; 46:925–928.
- 238. Fazio VW, Church JM. Complications and function of the continent ileostomy at the Cleveland Clinic. World J Surg. 1988;12:148–154.
- 239. Vernava AM 3rd, Goldberg SM. Is the Kock pouch still a viable option? *Int J Colorectal Dis.* 1988;3:135–138.
- 240. Fazio VW, Tjandra JJ. Technique for nipple valve fixation to prevent valve slippage in continent ileostomy. *Dis Colon Rectum*. 1992;35:1177–1179.
- 241. Ecker KW, Hildebrandt U, Haberer M, Feifel G. Biomechanical stabilization of the nipple valve in continent ileostomy. *Br J Surg.* 1996;83:1582–1585.
- 242. Litle VR, Barbour S, Schrock TR, Welton ML. The continent ileostomy: long-term durability and patient satisfaction. *J Gastrointest Surg.* 1999;3:625–632.
- 243. Wasmuth HH, Svinsås M, Tranø G, et al. Surgical load and long-term outcome for patients with Kock continent ileostomy. *Colorectal Dis.* 2007;9:713–717.
- 244. Ojerskog B, Hällström T, Kock NG, Myrvold HE. Quality of life in ileostomy patients before and after conversion to the continent ileostomy. *Int J Colorectal Dis.* 1988;3:166–170.
- 245. Parc R, Legrand M, Frileux P, Tiret E, Ratelle R. Comparative clinical results of ileal-pouch anal anastomosis and ileorectal anastomosis in ulcerative colitis. *Hepatogastroenterology*. 1989;36:235–239.
- 246. Khubchandani IT, Kontostolis SB. Outcome of ileorectal anastomosis in an inflammatory bowel disease surgery experience of three decades. *Arch Surg.* 1994;129:866–869.
- 247. Pastore RL, Wolff BG, Hodge D. Total abdominal colectomy and ileorectal anastomosis for inflammatory bowel disease. *Dis Colon Rectum.* 1977;40:1455–1464.
- 248. Saito Y, Sawada T, Tsuno N, et al. Total colectomy and ileorectal anastomosis in ulcerative colitis. *J Gastroenterol.* 1995; 8:131–134.
- 249. Mortier PE, Gambiez L, Karoui M, et al. Colectomy with ileorectal anastomosis preserves female fertility in ulcerative colitis. *Gastroenterol Clin Biol.* 2006;30:594–597.
- 250. Börjesson L, Lundstam U, Oresland T, Brevinge H, Hultén L. The place for colectomy and ileorectal anastomosis: a valid surgical option for ulcerative colitis? *Tech Coloproctol.* 2006;10:237–241.
- 251. Kvist N, Jacobsen O, Kvist HK, et al. Malignancy in ulcerative colitis. *Scand J Gastroenterol*. 1989;24:497–506.
- 252. Leijonmarck CE, Löfberg R, Ost A, Hellers G. Long-term results of ileorectal anastomosis in ulcerative colitis in Stockholm County. *Dis Colon Rectum.* 1990;33:195–200.
- 253. da Luz Moreira A, Kiran RP, Lavery I. Clinical outcomes of ileorectal anastomosis for ulcerative colitis. *Br J Surg.* 2010;97:65–69.
- 254. Lofberg R, Leijonmarck CE, Brostrom O. Mucosal dysplasia and DNA content in ulcerative colitis patients with ileorectal anastomosis: follow-up study in a definite patient group. *Dis Colon Rectum.* 1991; 34:566–571.
- 255. Grüner OP, Flatmark A, Naas R, Fretheim B, Gjone E. Ileorectal anastomosis in ulcerative colitis: results in 57 patients. *Scand J Gastroenterol*. 1975;10:641–646.
- 256. Paoluzi OA, Di Paolo MC, Ricci F, Pasquali C, Iacucci M, Paoluzi P. Ileo-rectal anastomosis in ulcerative colitis: results of a long-term follow-up study. *Ital J Gastroenterol*. 1994;26:392–397.

- 257. Herline AJ, Meisinger LL, Rusin LC, et al. Is routine pouch surveillance for dysplasia indicated for ileoanal pouches? *Dis Colon Rectum.* 2003;46:156–159.
- 258. Thompson-Fawcett MW, Marcus V, Redston M, Cohen Z, McLeod RS. Risk of dysplasia in long-term ileal pouches and pouches with chronic pouchitis. *Gastroenterology*. 2001;121:275–281.
- 259. Börjesson L, Willén R, Haboubi N, Duff SE, Hultén L. The risk of dysplasia and cancer in the ileal pouch mucosa after restorative proctocolectomy for ulcerative proctocolitis is low: a long-term term follow-up study. *Colorectal Dis.* 2004;6:494–498.
- 260. Coull DB, Lee FD, Anderson JH, McKee RF, Finlay IG, Dunlop MG. Long-term cancer risk of the anorectal cuff following restorative proctocolectomy assessed by p53 expression and cuff dysplasia. *Colorectal Dis.* 2007;9:321–327.
- Das P, Smith JJ, Lyons AP, Tekkis PP, Clark SK, Nicholls RJ. Assessment of the mucosa of the indefinitely diverted ileo-anal pouch. *Colorectal Dis.* 2008;10:512–517.
- 262. Hultén L, Willén R, Nilsson O, Safarani N, Haboubi N. Mucosal assessment for dysplasia and cancer in the ileal pouch mucosa in patients operated on for ulcerative colitis: a 30-year follow-up study. *Dis Colon Rectum.* 2002;45:448–452.
- 263. Church J. Ileoanal pouch neoplasia in familial adenomatous polyposis: an underestimated threat. *Dis Colon Rectum*. 2005;48:1708–1713.
- 264. Setti-Carraro P, Talbot IC, Nicholls RJ. Long-term appraisal of the histological appearances of the ileal reservoir mucosa after restorative proctocolectomy for ulcerative colitis. *Gut.* 1994; 35:1721–1777.
- 265. Nilubol N, Scherl E, Bub DS, et al. Mucosal dysplasia in ileal pelvic pouches after restorative proctocolectomy. *Dis Colon Rectum*. 2007;50:825–831.
- 266. Lavery IC, Sirimarco MT, Ziv Y, Fazio VW. Anal canal inflammation after ileal pouch-anal anastomosis: the need for treatment. *Dis Colon Rectum.* 1995;38:803–806.
- 267. Shen B, Lashner BA, Bennett AE, et al. Treatment of rectal cuff inflammation (cuffitis) in patients with ulcerative colitis following restorative proctocolectomy and ileal pouch-anal anastomosis. *Am J Gastroenterol*. 2004;99:1527–1531.
- 268. Shen B, Fazio VW, Remzi FH, et al. Risk factors for diseases of ileal pouch-anal anastomosis after restorative proctocolectomy for ulcerative colitis. *Clin Gastroenterol Hepatol.* 2006;4:81–89.
- 269. Sequens R. Cancer in the anal canal (transitional zone) after restorative proctocolectomy with stapled ileal pouch-anal anastomosis. *Int J Colorectal Dis.* 1997;12:254–255.
- 270. Rotholtz NA, Pikarsky AJ, Singh JJ, Wexner SD. Adenocarcinoma arising from along the rectal stump after double-stapled ileorectal J-pouch in a patient with ulcerative colitis: the need to perform a distal anastomosis. *Dis Colon Rectum*. 2001;44:1214–1217.
- 271. Baratsis S, Hadjidimitriou F, Christodoulou M, Lariou K. Adenocarcinoma in the anal canal following ileal pouch-anal anastomosis for ulcerative colitis using a double stapling technique. *Dis Colon Rectum.* 2002;45:687–692.
- 272. Lee SW, Sonoda T, Milsom JW. Three cases of adenocarcinoma following restorative proctocolectomy with hand-sewn anastomosis for ulcerative colitis: a review of reported cases in the literature. *Colorectal Dis.* 2005;7:591–597.
- 273. Branco BC, Sachar DB, Heimann TM, Sarpel U, Harpaz N, Greenstein AJ. Adenocarcinoma following ileal pouch-anal

- anastomosis for ulcerative colitis: review of 26 cases. *Inflamm Bowel Dis.* 2009;15:295–299.
- 274. Um JW, M'Koma AE. Pouch-related dysplasia and adenocarcinoma following restorative proctocolectomy for ulcerative colitis. *Tech Coloproctol.* 2011;15:7–16.
- 275. Panier-Suffat L, Marracino M, Resegotti A, et al. Anal transitional zone adenocarcinoma following restorative proctocolectomy for ulcerative colitis: case report and review of literature. *Acta Gastroenterol Belg.* 2009;72:441–443.
- 276. Candioli S, Manigrasso A, Arcieri S, et al. Adenocarcinoma following restorative proctocolectomy for ulcerative colitis: a case report and review of the literature [in Italian]. *G Chir.* 2007;28:371–376.
- 277. Chia CS, Chew MH, Chau YP, Eu KW, Ho KS. Adenocarcinoma of the anal transitional zone after double stapled ileal pouch-anal anastomosis for ulcerative colitis. *Colorectal Dis.* 2008;10:621–623.
- 278. Holder-Murray J, Fichera A. Anal transition zone in the surgical management of ulcerative colitis. *World J Gastroenterol*. 2009;15:769–773.
- 279. Lohmuller JL, Pemberton JH, Dozois RR, Ilstrup D, van Heerden J. Pouchitis and extraintestinal manifestations of inflammatory bowel disease after ileal pouch-anal anastomosis. *Ann Surg.* 1990;211:622–627.
- 280. Turina M, Pennington CJ, Kimberling J, Stromberg AJ, Petras RE, Galandiuk S. Chronic pouchitis after ileal pouch-anal anastomosis for ulcerative colitis: effect on quality of life. *J Gastrointest Surg.* 2006;10:600–606.
- 281. Penna C, Dozois R, Tremaine W, et al. Pouchitis after ileal pouch-anal anastomosis for ulcerative colitis occurs with increased frequency in patients with associated primary sclerosing cholangitis. *Gut.* 1996;38:234–239.
- 282. Abdelrazeq AS, Kandiyil N, Botterill ID, et al. Predictors for acute and chronic pouchitis following restorative proctocolectomy for ulcerative colitis. *Colorectal Dis.* 2008;10:805–813.
- 283. Gustavsson S, Weiland LH, Kelly KA. Relationship of backwash ileitis to ileal pouchitis after ileal pouch-anal anastomosis. *Dis Colon Rectum.* 1987;30:25–28.
- 284. Samarasekera DN, Stebbing JF, Kettlewell MG, Jewell DP, Mortensen NJ. Outcome of restorative proctocolectomy with ileal reservoir for ulcerative colitis: comparison of distal colitis with more proximal disease. *Gut.* 1996;38:574–577.
- 285. Schmidt CM, Lazenby AJ, Hendrickson RJ, Sitzmann JV. Preoperative terminal ileal and colonic resection histopathology predicts risk of pouchitis in patients after ileaanal pull-through procedure. *Ann Surg.* 1998;227:663–665.

- 286. Campieri M, Gionchetti P. Probiotics in inflammatory bowel disease: new insight to pathogenesis or a possible therapeutic alternative? *Gastroenterology*, 1999;116:1246–1249.
- 287. Mahadevan U, Sandborn WJ. Diagnosis and management of pouchitis. *Gastroenterology*. 2003;124:1636–1650.
- 288. Shen B, Achkar JP, Lashner BA, et al. Endoscopic and histologic evaluation together with symptom assessment are required to diagnose pouchitis. *Gastroenterology*. 2001;121:261–267.
- Fogt F, Deren JJ, Nusbaum M, Wellmann A, Ross HM. Pouchitis in ulcerative colitis: correlation between predictors from colectomy specimens and clinico-histological features. *Eur Surg Res*. 2006;38:407–413.
- 290. Shen B, Achkar JP, Connor JT, et al. Modified pouchitis disease activity index: a simplified approach to the diagnosis of pouchitis. *Dis Colon Rectum*. 2003;46:748–753.
- 291. Madden MV, McIntyre AS, Nicholls RJ. Double-blind crossover trial of metronidazole versus placebo in chronic unremitting pouchitis. *Dig Dis Sci.* 1994;39:1193–1196.
- 292. Sandborn WJ, Pardi DS. Clinical management of pouchitis. *Gastroenterology*. 2004;127:1809–1814.
- 293. Shen B, Achkar JP, Lashner BA, et al. A randomized clinical trial of ciprofloxacin and metronidazole to treat acute pouchitis. *Inflamm Bowel Dis.* 2001;7:301–305.
- 294. Gionchetti P, Rizzello F, Helwig U, et al. Prophylaxis of pouchitis onset with probiotic therapy: a double-blind, placebocontrolled trial. *Gastroenterology*. 2003;124:1202–1209.
- 295. Gionchetti P, Rizzello F, Venturi A, et al. Oral bacteriotherapy as maintenance treatment in patients with chronic pouchitis: a double-blind, placebo-controlled. *Gastroenterology*. 2000;119:305–309.
- 296. Holubar SD, Cima RR, Sandborn WJ, Pardi DS. Treatment and prevention of pouchitis after ileal pouch-anal anastomosis for chronic ulcerative colitis. *Cochrane Database Syst Rev.* 2010;(6):CD001176.
- 297. Sambuelli A, Boerr L, Negreira S, et al. Budesonide enema in pouchitis: a double-blind, double-dummy, controlled trial. *Aliment Pharmacol Ther.* 2002;16:27–34.
- 298. Bell AJ, Price AB, Forbes A, Ciclitira PJ, Groves C, Nicholls RJ. Pre-pouch ileitis: a disease of the ileum in ulcerative colitis after restorative proctocolectomy. *Colorectal Dis.* 2006;8:402–410.
- 299. McLaughlin SD, Clark SK, Bell AJ, Tekkis PP, Ciclitira PJ, Nicholls RJ. Incidence and short-term implications of prepouch ileitis following restorative proctocolectomy with ileal pouch-anal anastomosis for ulcerative colitis. *Dis Colon Rectum*. 2009;52:879–883.