

ONLINE FIRST

Increased Risk of Postoperative Deep Vein Thrombosis and Pulmonary Embolism in Patients With Inflammatory Bowel Disease

A Study of National Surgical Quality Improvement Program Patients

Andrea Merrill, MD; Frederick Millham, MD

Hypothesis: Patients with inflammatory bowel disease (IBD) undergoing surgery are at increased risk for postoperative thromboembolism, including deep vein thrombosis (DVT), pulmonary embolism (PE), myocardial infarction, and stroke.

Design: Retrospective cohort study.

Setting: Two hundred eleven hospitals participating in the American College of Surgeons National Surgical Quality Improvement Program.

Patients: All 271 368 patients from the National Surgical Quality Improvement Program 2008 Participant Use Data File were examined, and 2249 patients with IBD were compared with 269 119 patients without IBD.

Main Outcome Measures: Occurrence of DVT, PE, myocardial infarction, or stroke within 30 days of surgery.

Results: Of 268 703 National Surgical Quality Improvement Program patients, 2249 (0.8%) had IBD. There were 2665 cases of DVT or PE (1.0%). Occurrence of DVT or

PE was more common in patients with IBD (2.5%) overall ($P < .001$). Nonintestinal surgical cases had a higher rate of DVT or PE (5.0%; $P = .002$). Regression analysis, controlling for confounders, confirmed that IBD was associated with increased risk for DVT or PE (odds ratio = 2.03; 95% CI, 1.52-2.70). For nonintestinal surgery, risk of DVT or PE for patients with IBD was increased (odds ratio = 4.45; 95% CI, 1.72-11.49). Inflammatory bowel disease had no effect on risk of postoperative myocardial infarction or stroke.

Conclusions: Patients with IBD are at increased risk for developing postoperative DVT or PE. This risk persists when potential confounding variables are controlled for. Risk of DVT or PE appears to be even higher for patients with IBD who are having nonintestinal surgery. Cardiac and stroke risks do not appear to be increased by IBD. Perhaps standards for DVT and PE prophylaxis in these cases should be reconsidered.

Arch Surg. 2012;147(2):120-124. Published online October 17, 2011. doi:10.1001/archsurg.2011.297

PATIENTS WITH INFLAMMATORY bowel disease (IBD) are at increased risk for thromboembolic events, especially deep vein thrombosis (DVT) and pulmonary embolism (PE). This association was first described in 1936 by Barger and

gest that patients with IBD have increased risk of venous thromboembolism varying from as low as 1.5 times that of nonaffected patients^{2,3} to as high as 4.7-fold for patients with Crohn disease in a population-based cohort study.⁴ Moreover, patients with IBD who have PE appear to have increased risk of death compared with others.^{2,5} Despite these findings, standard DVT and PE prophylaxis guidelines do not recommend enhanced prophylaxis for patients with IBD.⁶⁻⁸ Recent data suggest that patients with IBD who require surgery are at very high risk for DVT. Scarpa et al⁹ have reported that, compared with patients with cancer undergoing colon resections, patients with ulcerative colitis have an odds ratio (OR) of 7.4 for having postoperative DVT.

We wondered whether patients with IBD undergoing surgery in the American College of Surgeons National Surgical Quality

 CME available online at www.jamaarchivescme.com and questions on page 105

See Invited Critique
at end of article

Author Affiliations:
Department of Surgery,
Massachusetts General Hospital
(Dr Merrill) and Department of
Surgery, Newton Wellesley
Hospital (Dr Millham), Boston.

Barker,¹ who described 18 patients with thromboembolic disease (predominantly venous) in more than 1000 patients treated for IBD at Mayo Clinic. Recent publications sug-

Table 1. Definition of Inflammatory Bowel Disease

ICD-9-CM Code in "Podiat" Field	Definition	Patients, No. % (n = 2249)
555	Regional enteritis	36 (1.6)
555.0	Regional enteritis of small intestine	284 (12.6)
555.1	Regional enteritis of large intestine	238 (10.6)
555.2	Regional enteritis of small intestine with large intestine	225 (10.0)
555.9	Regional enteritis, unspecified site	363 (16.1)
556	Ulcerative colitis	40 (1.8)
556.0	Ulcerative colitis, (chronic) enterocolitis	30 (1.3)
556.1	Ulcerative colitis, (chronic) ileocolitis	24 (1.1)
556.2	Ulcerative colitis, (chronic) proctitis	55 (2.4)
556.4	Ulcerative colitis, pseudopolyposis of colon	8 (0.4)
556.5	Ulcerative colitis, left-sided ulcerative colitis	47 (2.1)
556.6	Ulcerative colitis, universal colitis	244 (10.8)
556.8	Ulcerative colitis, other ulcerative colitis	101 (4.5)
556.9	Ulcerative colitis, unspecified	520 (23.1)

Abbreviation: ICD-9-CM, *International Classification of Diseases, Ninth Revision, Clinical Modification*.

Improvement Program (NSQIP) member hospitals had a similar risk of DVT or PE. The NSQIP collected data from more than 170 hospitals in 2008. These data were gathered by specifically trained nurses working from a rigorously defined and validated data definition dictionary. The purpose of the NSQIP is to provide risk-adjusted clinical outcomes data to participating hospitals. The NSQIP Participant Use Data File (PUF) is a deidentified research database available to NSQIP participants. As such, the NSQIP PUF data set presents an opportunity to examine the relationship of DVT and PE with IBD in a large group of patients for whom data on comorbid conditions and other potential confounding variables are available and well defined. Furthermore, hospitals participating in the NSQIP, having invested in quality improvement, might be expected to treat patients with best practices, at least with respect to DVT prophylaxis. This would therefore reduce the opportunity for treatment bias between centers. Because there may be a bias toward patients with complicated acute inflammatory conditions in the IBD group, we also wondered whether patients who have surgery on nonintestinal sites would have a risk of DVT or PE different from that of patients undergoing small- and/or large-bowel procedures.

METHODS

DATA ACQUISITION AND PATIENT SELECTION

We used the NSQIP PUF for 2008 to evaluate rates of DVT and PE in surgical patients with and without IBD. The NSQIP 2008 PUF contains complete records on 271 368 procedures or ad-

Table 2. Definition of Intestinal Surgery

CPT Codes	Definition	Patients, No. ^a (n = 84 539)
44005-44160	Incision or excision of small or large bowel	40 900
44180-44238	Laparoscopic procedure, small or large bowel	20 128
44300-44346	Formation of stomas	50
44602-44680	Closure, plication, or stricturoplasty	507
44900-44979	Surgery on appendix	22 465
45000-45999	Surgery on rectum	489

Abbreviation: CPT, *Current Procedural Terminology*.

^aA total of 2149 patients with inflammatory bowel disease had intestinal surgery.

missions. This database collects data from more than 200 participating hospitals around the country and includes 136 variables describing patient characteristics, historical features, process variables, and outcomes gathered retrospectively by specifically trained nurse reviewers. Permission to use this data set for research purposes was granted by the NSQIP. The Newton Wellesley Hospital Human Research Investigation Committee also approved of this work.

PREOPERATIVE VARIABLES

We dichotomized patient demographic characteristics, history, and preoperative laboratory data, taking care to indicate missing variables where appropriate. Crohn disease was defined as present in any case where the postoperative diagnosis field, "podiat," contained *International Classification of Diseases, Ninth Revision, Clinical Modification* (ICD-9-CM) codes 555-555.9 (regional enteritis). Ulcerative colitis was defined as present in any case where the podiat field contained ICD-9-CM codes 556-556.9 (ulcerative colitis). **Table 1** lists the ICD-9-CM codes defining the inclusion criteria for these diseases and the frequencies of each code in the database. Inflammatory bowel disease was defined as any case meeting criteria for either Crohn disease or ulcerative colitis. Intestinal surgery was defined as being present in any record with a principle procedure *Current Procedural Terminology* code indicating surgery on the small or large bowel. **Table 2** lists *Current Procedural Terminology* codes defining intestinal surgery.

OUTCOMES

The NSQIP variables are rigorously defined using a standardized dictionary. Data are abstracted by specifically trained nurses whose performance is subject to yearly validation. The NSQIP dictionary definitions of these occurrences are listed in the eAppendix (<http://www.archsurg.com>). We defined DVT as any record with a value greater than 0 recorded in the "nothdvt" field. Similarly, PE was defined as value greater than 0 in the "npul-embol" field, myocardial infarction (MI) as a value greater than 0 in the "ncdmi" field, and stroke as a value greater than 0 in the "ncnscva" field. Because DVT and PE are thought to represent different manifestations of the same process, the spontaneous formation of thrombus in the central veins and the migration of such thrombi centrally, we created a composite variable to identify any patient who had either DVT or PE. Similarly, because MI and cerebrovascular accident may represent spontaneous arterial thrombosis and migration of thrombi, we created a composite variable to identify patients who had either an MI or a stroke. Death was defined as having occurred in any

Table 3. Characteristics of National Surgical Quality Improvement Program Patients With and Without Inflammatory Bowel Disease

Characteristic	Patients, No. (%)		P Value ^a
	Non-IBD (n = 269 119)	IBD (n = 2249)	
DVT or PE	2608 (1.0)	57 (2.5)	<.001
Male	114 909 (42.7)	1122 (49.9)	<.001
Race/ethnicity			
White	206 511 (76.7)	1898 (84.4)	<.001
African American	26 915 (10.0)	133 (5.9)	<.001
Hispanic	2603 (1.0)	46 (2.0)	<.001
Admitted from home	260 178 (96.7)	2146 (95.4)	.002
Smoker	55 928 (20.8)	420 (18.7)	.01
BMI >30	104 119 (38.7)	444 (19.7)	<.001
Steroid use	7205 (2.7)	821 (36.5)	<.001
History			
Any dyspnea	30 353 (11.3)	122 (5.4)	<.001
Chemotherapy	2795 (1.0)	5 (0.2)	<.001
Metastatic cancer	5250 (2.0)	6 (0.3)	<.001
Brain tumor	515 (0.2)	3 (0.1)	.81
Weight loss	5920 (2.2)	214 (9.5)	<.001
Operation within last 30 d	7958 (3.0)	35 (1.6)	<.001
Outpatient surgery	92 822 (34.5)	55 (2.4)	<.001
Emergency surgery	34 176 (12.7)	147 (6.5)	<.001
ASA score of 1 or 2	147 296 (54.7)	1531 (68.1)	<.001
WBC count >12 000/ μ L	30 742 (11.4)	318 (14.1)	<.001
Postoperative organ space infection	3281 (1.2)	148 (6.6)	<.001
Postoperative sepsis	5155 (1.9)	140 (6.2)	<.001
Postoperative septic shock	3179 (1.2)	46 (2.0)	<.001
Dehiscence	1586 (0.6)	32 (1.4)	<.001
Failure to wean from ventilator	6141 (2.3)	60 (2.7)	.23
Postoperative reintubation	3644 (1.4)	33 (1.5)	.65
Postoperative neurologic deficit	175 (0.1)	3 (0.1)	.18
Postoperative CVA or MI	1164 (0.4)	9 (0.4)	.68
Postoperative cardiac arrest	1156 (0.4)	7 (0.3)	.52
Postoperative urinary tract infection	4329 (1.6)	77 (3.4)	<.001
Death	4608 (1.7)	26 (1.2)	.04
Age, mean (SEM), y	55.5 (0.03)	43.3 (0.33)	<.001
Total work RVUs, mean (SEM)	16.0 (0.02)	25.9 (0.17)	<.001
Total LOS, mean (SEM), d	4.4 (0.02)	9.1 (0.21)	<.001

Abbreviations: ASA, American Society of Anesthesiologists; BMI, body mass index (calculated as weight in kilograms divided by height in meters squared); CVA, cerebrovascular accident; DVT, deep vein thrombosis; IBD, inflammatory bowel disease; LOS, length of stay; MI, myocardial infarction; PE, pulmonary embolism; RVUs, relative value units; WBC, white blood cell.

SI conversion factor: To convert WBC count to $\times 10^9$ per liter, multiply by 0.001.

^aThe *P* values for categorical variables are from Fisher exact test; the *P* values for continuous variables are from 2-sided *t* test.

patient whose record had the date 2008 in the “yrdeath field.” By NSQIP definition, this designation includes patients who died in the hospital or within 30 days of surgery.

STATISTICAL ANALYSIS

We tested for variables associated with DVT or PE among all of the variables in the NSQIP data set using univariate methods: χ^2 test for categorical variables and *t* test for continuous variables. These findings then informed the construction of a number of logistic regression models. Final models were those with the most

parsimonious use of covariates and the greatest explanatory power. All analysis was performed in Stata version 10.1 statistical software (StataCorp LP, College Station, Texas).

RESULTS

The NSQIP 2008 PUF database included 271 368 patients who underwent surgery. Characteristics of patients with and without IBD are shown in **Table 3**. Patients with and without IBD differed in many ways. There were 2249 patients (0.8%) identified as having a diagnosis of IBD, 57 of whom (2.5%) had DVT or PE, compared with 1.0% of patients without IBD ($P < .001$). A total of 1164 patients (0.4%) without IBD and 9 patients (0.4%) with IBD had a cerebrovascular accident or MI ($P = .68$). Patients with IBD were more commonly male and white or Hispanic than patients without IBD ($P < .001$). Patients with IBD were younger, had on average more relative value units per procedure, and had a longer average length of stay. Patients with IBD were less commonly admitted from home, were smokers, or were obese ($P < .001$). More than 36% of patients with IBD had used steroids compared with fewer than 3% of patients without a diagnosis of IBD ($P < .001$). Patients with IBD were less likely to undergo surgery as outpatients or emergencies ($P < .001$) and more likely to have American Society of Anesthesiologists scores of 1 or 2 ($P < .001$), but they were more likely to have an elevated white blood cell count, have an organ space infection, sepsis, or septic shock, or to dehiscence their fascial closure ($P < .001$). However, they had lower postoperative mortality rates ($P = .04$).

Of the multivariate logistic regression models constructed to try to account for these various confounding factors, the most efficient is shown in **Table 4**. This model includes all variables available in the NSQIP that add to the power to predict DVT or PE. When these numerous confounders are controlled for, IBD remains a significant predictor of DVT or PE (OR = 2.03; 95% CI, 1.52-2.70). In fact, only a history of brain tumor or postoperative neurologic deficit have a higher OR in this model. Mortality occurred in 224 of 2608 patients without IBD who had DVT or PE (8.6%) and 5 of 57 patients without IBD who had DVT or PE (8.8%) ($P > .99$).

When patients with procedures on the small and large bowel were excluded, 100 patients with IBD remained. Of these, 5 (5.0%) had DVT or PE compared with 1778 cases of DVT or PE among 213 402 patients without IBD (0.8%) ($P = .002$). When a regression model is constructed for this subset of patients undergoing nonintestinal procedures, the patients with IBD are found to have a 4.45-fold increased risk of DVT or PE ($P = .002$) compared with patients without IBD.

We repeated the modeling exercise for the composite outcome of perioperative MI and stroke. This analysis revealed no association between IBD and this outcome or its individual components.

COMMENT

An increased risk of DVT and PE in patients with IBD has been evident for the past 75 years. Most work in this area has not looked specifically at patients undergoing

surgery. Patients with IBD frequently require surgical intervention, and an understanding of their risk of venous thromboembolism is therefore an important issue. This study aimed to examine rates of DVT and PE in patients with IBD undergoing surgery using data from the NSQIP.

Our findings are consistent with others in this area. We found that IBD increases the risk of perioperative DVT or PE 2-fold (OR=2.03; 95% CI, 1.52-2.70). Bernstein and Nabalamba³ found that in patients hospitalized in Canada, IBD resulted in a DVT relative risk of 1.32 to 1.53, depending on age group. Using a survey method, Miehsler et al¹⁰ found that 6.2% of patients with IBD reported an episode of thromboembolism, resulting in an OR of 3.6 (95% CI, 1.7-7.8). Most recently, using data from the National Inpatient Sample, Nguyen and Sam¹¹ found that hospitalized patients with IBD had an OR for DVT or PE of 2.5 (95% CI, 1.83-3.43).

The only previous study to examine perioperative DVT, a report of the 7-year experience of a single institution where patients with ulcerative colitis and colorectal cancer both received low-molecular-weight heparin at a dosage of 4000 IU/d, found that patients with ulcerative colitis had an OR of 7.4 (95% CI, 1.14-44.4) for experiencing DVT.⁹ The broad CIs in this relatively small study suggest that the statistical model may be unstable. Yet, the result comports well with our findings.

We find no support for the claim that DVT or PE in the setting of IBD is more lethal than DVT or PE occurring in the absence of IBD as has been described by Nguyen and Sam¹¹ and Murthy and Nguyen.²

Arterial thromboembolic events are also reported to be increased in patients with IBD.¹² We found no support for this conclusion in our data. In this case, we may have been limited by the lack of data on arterial thrombotic events not involving the coronary or cerebral vessels. It may be that if arterial thromboembolism were a reported NSQIP complication, such a relationship would appear.

This increased risk of DVT or PE in patients with IBD has been shown to be an extraintestinal manifestation specific to IBD and not solely due to the inflammatory state of the disease. Miehsler et al¹⁰ compared the risk of thromboembolism in patients with IBD, celiac disease, and rheumatoid arthritis and control subjects. While they found an association between increased frequency of thromboembolic events and IBD, there was no such association in patients with the inflammatory conditions of celiac disease or rheumatoid arthritis.

Many theories have been proposed to explain this extraintestinal manifestation of IBD. Factors implicated in the pathogenesis of thromboembolism in patients with IBD include platelet hyperactivation mediated by the CD40-CD40 ligand pathway, hyperhomocysteinemia (secondary to malabsorption and medications), increased coagulation factors, and impaired fibrinolysis.¹³⁻¹⁷ Furthermore, studies have even shown a paradoxical improvement in symptoms of inpatients with ulcerative colitis treated with heparin.^{18,19}

There are important strengths and weaknesses associated with using the NSQIP PUF data set for outcomes research. Among the weaknesses, perhaps the most important is that the NSQIP was designed to compare overall

Table 4. Factors Associated With Deep Vein Thrombosis or Pulmonary Embolism in the National Surgical Quality Improvement Program^a

Factor	OR (95% CI)	P Value
Demographic variable		
Male	1.14 (1.05-1.23)	.002
African American	1.09 (0.96-1.24)	.17
Hispanic	0.75 (0.61-0.93)	.007
Admitted from home	0.79 (0.69-0.90)	<.001
Smoker	0.86 (0.78-0.95)	.004
BMI>30	1.19 (1.09-1.29)	<.001
Age ^b	1.02 (1.01-1.02)	<.001
Medical history		
Brain tumor	3.06 (2.07-4.52)	<.001
Steroid use	1.38 (1.18-1.60)	<.001
Any dyspnea	1.11 (1.01-1.23)	.03
Chemotherapy	1.81 (1.42-2.31)	<.001
Metastatic cancer	1.33 (1.10-1.60)	.003
Weight loss	1.35 (1.15-1.59)	<.001
Operation within last 30 d	1.28 (1.10-1.48)	.001
Clinical factor		
IBD	2.03 (1.52-2.70)	<.001
Outpatient procedure	0.29 (0.24-0.34)	<.001
Total LOS ^b	1.01 (1.01-1.02)	<.001
Emergency surgery	1.26 (1.13-1.40)	<.001
Total work RVUs ^b	1.02 (1.02-1.02)	<.001
ASA score of 1 or 2	0.76 (0.68-0.84)	<.001
WBC count >12 000/ μ L	1.24 (1.12-1.38)	<.001
Postoperative organ space infection	1.52 (1.28-1.80)	<.001
Postoperative sepsis	2.04 (1.78-2.34)	<.001
Postoperative septic shock	1.58 (1.35-1.86)	<.001
Dehiscence	1.42 (1.12-1.80)	.004
Failure to wean from ventilator	1.91 (1.66-2.20)	<.001
Postoperative reintubation	1.57 (1.34-1.84)	<.001
Postoperative neurologic deficit	2.71 (1.27-5.81)	.01
Postoperative CVA	1.98 (1.47-2.66)	<.001
Postoperative MI	1.51 (0.99-2.30)	.05
Postoperative cardiac arrest	1.40 (1.10-1.79)	.007
Postoperative urinary tract infection	1.60 (1.36-1.88)	<.001

Abbreviations: ASA, American Society of Anesthesiologists; BMI, body mass index (calculated as weight in kilograms divided by height in meters squared); CVA, cerebrovascular accident; IBD, inflammatory bowel disease; LOS, length of stay; MI, myocardial infarction; OR, odds ratio; RVUs, relative value units; WBC, white blood cell.

SI conversion factor: To convert WBC count to $\times 10^9$ per liter, multiply by 0.001.

^aModel $R^2 = 0.1426$.

^bContinuous variables.

outcomes across many hospitals. It was not designed to answer specific research questions regarding specific diseases or procedures. The NSQIP is a database of operations where postoperative diagnosis is cataloged, but most other diagnoses are not specifically recorded. In this study, it is possible that some patients with IBD for whom this was not the principle postoperative diagnosis may have been missed, therefore introducing a classification bias.

Second, because this is a database of operations and not of individual patients, it is possible that particular patients who had multiple admissions during 2008 may be captured more than once in the data set. Presumably the size of the data set would minimize the effect of multiple admissions, but where IBD was a relatively rare diagnosis and the fact that it often requires multiple hos-

pitalizations may create an opportunity for sample bias toward sicker patients with IBD.

There are a number of strengths in the NSQIP that we believe outweigh these potential sources of bias. First, these data are gathered by specifically trained nurses, accountable to a rigid quality assurance program and working from a well-defined data dictionary. To our knowledge, no other data set offers such reliable and well-validated information. For an investigation like this one, aiming to test an association between 2 relatively uncommon problems, only a prospective multicenter trial enrolling thousands of patients could approach this question. The NSQIP provides an economical alternative.

The second advantage to using the NSQIP is that it not only aggregates data from a large number of centers but also biases selection toward centers committed to quality improvement. One may assume that most, if not all, of the contributing centers are aware of best practices (Surgical Care Improvement Program, for example) and engage in state-of-the-art DVT and PE prophylaxis and detection. No other administrative data set has a bias toward quality that the NSQIP has.

In conclusion, this study of patients enrolled in the NSQIP database demonstrates that patients with IBD who undergo surgery have a 2-fold increased risk of DVT or PE. In patients with IBD who are having nonintestinal surgery, this risk may be even higher. These findings suggest that standard DVT and PE prophylaxis should be reconsidered for this patient group.

Accepted for Publication: July 22, 2011.

Published Online: October 17, 2011. doi:10.1001/archsurg.2011.297

Correspondence: Frederick Millham, MD, Department of Surgery, Newton Wellesley Hospital, 2000 Washington St, Ste 670, Green MOB, Newton, MA 02462 (fmillham@partners.org).

Author Contributions: Study concept and design: Merrill and Millham. Analysis and interpretation of data: Millham. Drafting of the manuscript: Merrill and Millham. Critical revision of the manuscript for important intellectual content: Merrill and Millham. Statistical analysis: Millham.

Financial Disclosure: None reported.

Online-Only Material: The eAppendix is available at <http://www.archsurg.com>.

REFERENCES

1. Barga JA, Barker NW. Extensive arterial and venous thrombosis complicating chronic ulcerative colitis. *Arch Intern Med*. 1936;58(1):17-31.
2. Murthy SK, Nguyen GC. Venous thromboembolism in inflammatory bowel disease: an epidemiological review. *Am J Gastroenterol*. 2011;106(4):713-718.
3. Bernstein CN, Nabalamba A. Hospitalization-based major comorbidity of inflammatory bowel disease in Canada. *Can J Gastroenterol*. 2007;21(8):507-511.
4. Bernstein CN, Blanchard JF, Houston DS, Wajda A. The incidence of deep venous thrombosis and pulmonary embolism among patients with inflammatory bowel disease: a population-based cohort study. *Thromb Haemost*. 2001;85(3):430-434.
5. Solem CA, Loftus EV, Tremaine WJ, Sandborn WJ. Venous thromboembolism in inflammatory bowel disease. *Am J Gastroenterol*. 2004;99(1):97-101.
6. Carter MJ, Lobo AJ, Travis SP; IBD Section, British Society of Gastroenterology. Guidelines for the management of inflammatory bowel disease in adults. *Gut*. 2004;53(suppl 5):V1-V16.
7. Authors T; Thromboembolic Risk Factors (THRIFT) Consensus Group. Risk of and prophylaxis for venous thromboembolism in hospital patients. *BMJ*. 1992;305(6853):567-574.
8. Verstraete M. Prophylaxis of venous thromboembolism. *BMJ*. 1997;314(7074):123-125.
9. Scarpa M, Pilon F, Pengo V, et al. Deep venous thrombosis after surgery for inflammatory bowel disease: is standard dose low molecular weight heparin prophylaxis enough? *World J Surg*. 2010;34(7):1629-1636.
10. Miehsler W, Reinisch W, Valic E, et al. Is inflammatory bowel disease an independent and disease specific risk factor for thromboembolism? *Gut*. 2004;53(4):542-548.
11. Nguyen GC, Sam J. Rising prevalence of venous thromboembolism and its impact on mortality among hospitalized inflammatory bowel disease patients. *Am J Gastroenterol*. 2008;103(9):2272-2280.
12. Bernstein CN, Wajda A, Blanchard JF. The incidence of arterial thromboembolic diseases in inflammatory bowel disease: a population-based study. *Clin Gastroenterol Hepatol*. 2008;6(1):41-45.
13. Irving PM, Pasi KJ, Rampton DS. Thrombosis and inflammatory bowel disease. *Clin Gastroenterol Hepatol*. 2005;3(7):617-628.
14. Twig G, Zandman-Goddard G, Szyper-Kravitz M, Shoenfeld Y. Systemic thromboembolism in inflammatory bowel disease: mechanisms and clinical applications. *Ann N Y Acad Sci*. 2005;1051:166-173.
15. Danese S, Katz JA, Saibeni S, et al. Activated platelets are the source of elevated levels of soluble CD40 ligand in the circulation of inflammatory bowel disease patients. *Gut*. 2003;52(10):1435-1441.
16. Webberley MJ, Hart MT, Melikian V. Thromboembolism in inflammatory bowel disease: role of platelets. *Gut*. 1993;34(2):247-251.
17. Cattaneo M, Vecchi M, Zighetti ML, et al. High prevalence of hyperhomocysteinemia in patients with inflammatory bowel disease: a pathogenic link with thromboembolic complications? *Thromb Haemost*. 1998;80(4):542-545.
18. Folwaczny C, Wiebecke B, Loeschke K. Unfractionated heparin in the therapy of patients with highly active inflammatory bowel disease. *Am J Gastroenterol*. 1999;94(6):1551-1555.
19. Gaffney PR, Doyle GT, Gaffney A, Hogan J, Hayes DP, Annis P. Paradoxical response to heparin in 10 patients with ulcerative colitis. *Am J Gastroenterol*. 1995;90(2):220-223.

INVITED CRITIQUE

Use of National Surgical Quality Improvement Program Outcomes Data to Change Guidelines

A Sound Idea?

Merrill and Millham¹ provide compelling data to associate patients with inflammatory bowel disease (IBD) with an increased postoperative risk for deep venous thrombosis (DVT) and pulmonary embolism. Using National Surgical Quality

Improvement Program (NSQIP) data, the results likely confirm what many physicians who treat patients with IBD already believe. Current guidelines from the American College of Chest Physicians 2008 Conference on Antithrombotic and Thrombolytic Therapy