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Biological Treatment and the Potential Risk of Adverse Postoperative Outcome in Patients With Inflammatory Bowel Disease: An Open-Source Expert Panel Review of the Current Literature and Future Perspectives

Open Source Research Collaborating Group (#OpenSourceResearch)*

Background: There is widespread concern that treatment with biologic agents may be associated with suboptimal postoperative outcome after surgery for inflammatory bowel diseases (IBD).

Aim: We aimed to search and analyze the literature regarding the potential association of biologic treatment on adverse postoperative outcome in patients with IBD. We used the subject as a case in point for surgical research. The aim was not to conduct a new systematic review.

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Method: This is an updated narrative review written in a collaborative method by authors invited through Twitter via the following hashtags (#OpenSourceResearch and #SoMe4Surgery). The manuscript was presented as slides on Twitter to allow discussion of each section of the paper sequentially. A Google document was created, which was shared across social media, and comments and edits were verified by the primary author to ensure accuracy and consistency.

Results: Forty-one collaborators responded to the invitation, and a total of 106 studies were identified that investigated the potential association of preoperative biological treatment on postoperative outcome in patients with IBD. Most of these studies were retrospective observational cohorts: 3 were prospective, 4 experimental, and 3 population-based studies. These studies were previously analyzed in 10 systematic/narrative reviews and 14 meta-analyses. Type of biologic agents, dose, drug concentration, antidrug antibodies, interval between last dose, and types of surgery varied widely among the studies. Adjustment for confounders and bias control ranged from good to very poor. Only 10 studies reported postoperative outcome according to Clavien–Dindo classification.

Conclusion: Although a large number of studies investigated the potential effect of biological treatment on postoperative outcomes, many reported divergent results. There is a need for randomized controlled trials. Future studies should focus on the avoiding the weakness of prior studies we identified. Seeking collaborators and sharing information via Twitter was integral to widening the contributors/authors and peer review for this article and was an effective method of collaboration.

Key Words: Crohn disease, inflammatory bowel disease, ulcerative colitis, biologic treatment, biologics, anti-TNF alpha, postoperative outcome, surgery.

WHAT DOES THIS ARTICLE ADD TO EXISTING LITERATURE AND WHAT IT DOES NOT ADD?

This study represents the most extensive literature review done on the subject using a unique, social media-based, methodology. It identified all the studies published until September 30, 2018, analyzed these studies, described the 106 studies, and attempted to answer these questions:

1. Why these 106 studies reported divergent results?
2. Why there are a large number of studies with severe limitations in methodology? Is this not a waste of resources?
3. Can social media and internet connection improve research quality and facilitate optimal design of future studies?

To the best of our knowledge, this represents the first research project conducted using social media to recruit coauthors and paragraph-by-paragraph revision of the manuscript via the #OpenSourceResearch.

The authors would like to emphasize that:

1. This is not a systematic review or meta-analysis because there are already many systematic reviews and meta-analysis about the subject with divergent conclusions.
2. This is a case in point for surgical research. With such study, the authors hope that less temptation to conduct small series studies will be encountered and better cooperation in conducting larger-scale studies will be encouraged to advance surgical science.

INTRODUCTION

Biologic therapies have revolutionized the management of inflammatory bowel diseases (IBD), that is Crohn disease (CD) and ulcerative colitis (UC). Biologics, including antitumor necrosis factor-alpha agents (a-TNF), and more recently anti-integrin and anti-IL-12/23 agents, are reserved for patients with moderate-to-severe disease activity or for patients who are medically refractory to conventional therapy such as azathioprine, mercaptopurine, methotrexate, or corticosteroids. However,

TNF is an important component of the immune defense mechanism and plays a role in wound healing through a dose-dependent effect on angiogenesis¹ and collagen synthesis.²⁻⁴ Inhibition of TNF-mediated pathways may impair wound healing after surgery, thus theoretically increasing the risk of postoperative complications such as surgical site infection and anastomosis-related complications, although the latter has, to date, never been demonstrated in any clinical study or trial.

Despite having been used since 1998, the natural history of CD appears to be unaffected using a-TNF.⁵ Hence, the risk of surgery to treat refractory CD may not have changed in the era of biologics. The overall risk of surgery was 22% in a recent European study,⁶ and risk for second surgery is 28.7% based on meta-analysis of population-based studies.⁷ Furthermore, up to 50% of CD patients have been exposed to an a-TNF at time of their first surgery.⁸ In UC, the risk of colectomy seems to have decreased in the era of biologics.⁹ It is not clear whether this change is related to the introduction of biologics or better preoperative optimization.

To date, there are 106 scientific papers assessing the effect of a-TNF therapy on postoperative outcome, with divergent conclusions. As such, there is a need to assess the current evidence and to assess each studies strength and weakness, so as to help plan future studies with optimal design and methodology. Thus, the aim of this study was to assess the potential association between preoperative exposure to biologics with postoperative outcomes in IBD patients and to make recommendations for the optimal design of future studies.

METHOD

This study is a narrative review based on a published systematic review by one of the authors (A.E.).¹⁰ In this update, all eligible studies were included. No statistical analyses or bias control analyses were conducted because of the wide heterogeneity of the included studies. Biologic treatment was defined as treatment with a-TNF agents (eg, infliximab or adalimumab), integrin inhibitors (eg, vedolizumab), or IL-12/23 inhibitors (eg, ustekinumab).

Aim

The aim of this review is to provide a case in point about surgical research by examining one subject in mintious details to identify the limitations of reported studies and attempt to lead the design of future studies.

Eligibility Criteria

Case-control and cohort studies were included irrespective of publication status, year of publication, or language. Included studies assessed patients with CD or UC undergoing laparoscopic or open abdominopelvic surgery. Based on pharmacokinetic studies, the intervention group included patients who received any type or dose of biologics within 3 months of surgery.

Outcome Measures

Outcome measures were assessed after 30 days of follow-up. The outcomes were assessed as defined by the authors of the included studies.

Search Strategy and Method of Updating the Review

The search strategy is attached as a [Supplementary Material](#). The search was prospectively updated by the authors using:

1. Alerts from PubMed.gov.
2. Alerts from researchgate.com (any citation of articles by authors).
3. Alerts from relevant journals.
4. Attending relevant conferences (European Society of ColoProctology [ESCP], European Crohn's and Colitis Organization [ECCO], Association of Coloproctology of Great Britain and Ireland, American Society of Colon and Rectal Surgeons [ASCRS]).
5. Following topical developments in the subject as the authors are reviewers in many international journals and are, or have been, members of guidelines committees for UC and CD with the following organization: European Crohn's and Colitis Organization, Crohn's and Colitis Foundation, and the American Society of Colon and Rectal Surgeons.
6. Contacting experts in the field using the Twitter #SoMe4Surgery hashtag. Many authors are expert in this subject (A.H., S.H., P.K., P.M., J.D., A.E., A.S., N.Y., S.W., see [Supplementary Material](#) with list of all authors). All the contributors were asked to update the list of the included articles using their Twitter network to ensure that all relevant studies are included. Two additional papers were identified via this mechanism.

Open Source Research Project

The paper was written in a collaborative method with authors invited personally and through Twitter via the following hashtags (#OpenSourceResearch and #SoMe4Surgery). A Google document was created which was shared across social media and comments and edits were verified by the primary author to ensure accuracy and consistency. The manuscript was presented on Twitter in sequential posts by section. Each post

contained 1–2 paragraphs of the manuscript modified to fit the limited space in Twitter and include powerpoint slides and images. This collaboration allowed this research and paper to be unique in the way it was written and edited.

Twitter offered a platform to engage researchers, to broaden the search, and to conduct scientific discussions.

The final draft was sent by email to all contributors for feedback prior to submission and the primary author (A.E.) completed the final draft which was then edited for grammar by a native English-speaking author (S.H.).

Twitter Analysis of the #OpenSourceResearch

Twitter is an American-based yet international online news and social networking service to which users post and interact with messages known as “tweets.” Tweets were originally restricted to 140 characters, but this limit was doubled for all languages except Chinese, Japanese, and Korean. Registered users may post, like, and “retweet” tweets, but unregistered users can only read them. Users access Twitter through its website interface, short message service, or its mobile-device application software (“app”).

As this paper was written as a social media-based collaboration, it was important to capture social media activity as potential source material for the paper and to assess the response to the idea of an open-source research paper, as well as to document the main influencers of these discussions. Data were collected through 2 tools—“Followthehashtag” for geographical mapping and gender split of tweeters, and “NodeXL” to document the networking interactions between tweeters and describe the interactions between these tweeters (replies, retweets, and mentions of other tweeters). Both tools can be used to quantify the number of tweeters, retweeters, and tweets. Both tools also provide information about individual tweets. Additional information is available at <http://analytics.followthehashtag.com/#/?id=dashboard> and at <https://www.smrfoundation.org/nodexl/>.

RESULTS

We identified a total of 106 studies that investigated the impact of preoperative biological treatment on postoperative outcome in patients with IBD. The relation of a-TNF therapy with postoperative outcomes in patients with IBD has been investigated in 32 retrospective CD cohorts^{11–42} (CD with/without UC), in 16 retrospective cohorts (UC),^{43–58} 3 prospective ones,^{59–61} 4 experimental studies,^{62–65} 3 population-based studies,^{66–68} 10 narrative reviews,^{69–78} and 14 meta-analyses^{10, 79–89} making a total of 82 studies over the past 15 years ([Tables 1–4](#)). In addition, 27 studies investigated this relation as part of other risk factors of unfavorable postoperative outcome, as well as studies with a focus on rheumatoid arthritis ([Table 5](#)). The 54 clinical studies, which included 32 retrospective cohorts of CD, 16 retrospective cohorts of UC, 3 prospective cohorts of CD, and 3 nation-based studies, are demonstrated in details in [Tables 1–3](#). In total, these clinical studies have included 22,923 patients of whom 5501 (24%) were on preoperative biological treatment.

TABLE 1. Clinical Studies About Type of Preoperative Biological Treatment in Patients With Crohn Disease

Author/Publication's Year	Cohort/Treated (7895/2026)	Type of Biological Treatment	Weeks Before Surgery
Observational retrospective studies			
1 Tay ¹¹ et al (2003)	100/22	IFX	8
2 Colombel ¹² et al (2004)	270/52	IFX	8 ^a
3 Marchal ²³ et al (2004)	79/40	IFX	12 ^b (8, 4, and less than 4 wk)
4 Appau ³⁴ et al (2008)	389/60	IFX	12
5 Indar ³⁷ et al (2009)	112/17	Anti-TNF	8
6 Nasir ³⁸ et al (2010)	370/119	IFX, ADA, CZM	8 ^a
7 Kasperek ³⁹ et al (2011)	94/48	IFX	12
8 Canedo ⁴⁰ et al (2011)	225/65	IFX	8
9 Regueiro ⁴¹ et al (2011)	24/11	IFX	2–4 wk after surgery ^c
10 El-Hussuna ⁴² et al (2012)	417/32	IFX, ADA, CZM	12
11 White ¹³ et al (2012)	338/59	IFX, ADA, CZM	12
12 Myreli ¹⁴ et al (2013)	298/111	IFX, ADA	8
13 Serradori ¹⁵ et al (2013)	217/42	Anti-TNF	12 (and 8 wk)
14 Syed ¹⁶ et al (2013)	325/150	Anti-TNF	8
15 Uchino ³⁶ et al (2013)	405/79	IFX	12
16 Bafford ¹⁷ et al (2013)	196/35	Anti-TNF	12
17 Kotze ¹⁸ et al (2016)	123/71	IFX, ADA	8
18 Shim ²¹ et al (2016)	60/60	Anti-TNF, UST	Up to 72
19 Zimmerman ²² et al (2016)	123/24 ^d	IFX	12 (8 and 4 wk)
20 Kotze ⁹⁷ et al (2017)	123/71	ADA	2
21 Jouvin ¹⁹ et al (2018)	360/58	IFX, ADA, CZM	8
22 Lightner ³² et al (2018)	213/213 (169 anti-TNF- α)	Anti-TNF- α and UST	12
Prospective studies			
23 Brouquet ⁵⁹ et al (2016)	592/340	IFX, ADA, VDZ, other anti-TNF	12
24 Fumery ⁶⁰ et al (2016)	209/93	IFX, ADA	12 (and 4 wk)
Nation-wide database study			
25 Nørgård ⁶⁶ et al (2013)	2293/214	IFX, ADA, CZM	12 (2 and 4 wk)

The time interval between last administered dose of biological treatment and surgery varied among the studies.

^aAnd 4 wk after surgery.

^bNine patients received anti-TNF- α more than 12 wk prior to surgery.

^cData were collected from a randomized clinical trial to investigate postoperative recurrence of CD.

^dPediatric patients.

IFX, infliximab; ADA, adalimumab; CZM, certolizumab; UST, ustekinumab; VDZ, vedolizumab; anti-TNF, antitumor necrosis factor agents.

TABLE 2. Clinical Studies About Type of Preoperative Biological Treatment in Patients With Crohn Disease, Ulcerative Colitis, and Indeterminate Colitis (Mixed Population Studies)

Author/Publication's Year		Cohort/Treated (3003/1234)	Type of Biological Treatment	Weeks Before Surgery
Observational retrospective studies				
1	Kunitake ²⁵ et al (2008)	413/101	IFX	12
2	Regadas ²⁶ et al (2010)	249/28	IFX	8
3	Rizzo ²⁷ et al (2011)	114/54	Anti-TNF	12 (and 4 wk)
4	Krane ²⁸ et al (2013)	518/142	IFX	12
5	Waterman ²⁹ et al (2013)	282/73	IFX, ADA	25
6	Lau ³⁰ et al (2015)	217/143	IFX, ADA, CZM	Not specified ^a
7	Alsaleh ³¹ et al (2016)	47/47	IFX	12
8	Lightner ³² et al (2016)	392/220	VDZ	12
9	Shwaartz ³³ et al (2016)	282/73	IFX, ADA, CZM	8
10	Yamada ³⁵ et al (2017)	443/193	IFX, ADA, VDZ	4
Prospective studies				
11	El-Hussuna ⁶¹ et al (2018)	46/18	IFX, ADA, CZM	12

^aAbout 65% of patients in this cohort received anti-TNF- α therapy before surgery.

IFX, infliximab; ADA, adalimumab; CZM, cemzia; VDZ, vedolizumab; anti-TNF, antitumor necrosis factor agents.

TABLE 3. Clinical Studies About Type of Preoperative Biological Treatment in Patients With Ulcerative Colitis

Author/Publication's Year		Cohort/Treated (11,965/2181)	Type of Biological Treatment	Weeks Before Surgery
Observational retrospective studies				
1	Selvasekar ⁴³ et al (2007)	301/47	IFX	8 ^a
2	Schluender ⁴⁴ et al (2007)	151/17	IFX	Up to 54
3	Mor ⁵¹ et al (2008)	523/85	IFX	Up to 37
4	Ferrante ⁵² et al (2009)	141/22	IFX	Up to 12
5	Coquet-Reinier ⁵³ et al (2010)	26/13	IFX	Up to 23
6	Gainsbury ⁵⁴ et al (2011)	81/29	IFX	12
7	Bregnbak ⁵⁵ et al (2012)	71/20	IFX	12
8	Kennedy ⁹⁸ et al (2012) ^b	38/11	IFX	8
9	Uchino ⁵⁷ et al (2013)	196/22	IFX	12
10	Eshuis ⁵⁸ et al (2013)	72/38	IFX	Up to 32
11	Gu ⁴⁵ et al (2013)	181/25	IFX, ADA, CZM	12
12	Nelson ⁴⁶ et al (2014)	78/28	IFX	1
13	Zittan ⁴⁷ et al (2016)	562/196	Anti-TNF	Up to 24
14	Kulaylat ⁴⁸ et al (2017)	2476/650	IFX, ADA, CZM	12
15	Lightner ⁴⁹ et al (2017)	150/150	VDZ	12
16	Ferrante ⁵⁰ et al (2017)	170/94	VDZ, anti-TNF	8–16
Nation-wide database study				
17	Nørgård ⁶⁷ et al (2013)	1226/199	Anti-TNF	12
18	Ward ⁶⁸ et al (2017)	6225/753	Anti-TNF	12 (and 4 wk)

^aOnly 49% of patients in the study cohort.

^bPediatric age cohort.

IFX, infliximab; ADA, adalimumab; CZM, cemzia; VDZ, vedolizumab; anti-TNF, antitumor necrosis factor agents.

How Confounding Factors Were Addressed in the Different Studies?

The included studies varied in the method in which they addressed potential confounding factors as type of medication, time interval between medication and surgery, drug concentration, preexistence of antidrug antibodies. Here is an account of these confounding factors and how they were addressed in different studies.

Type of medication

Some studies defined the type of biological treatments (e.g. infliximab); others reported a broad category of treatment (e.g. a-TNF, including certolizumab pegol and golimumab) while a few studies included all biological treatments without any definition (Tables 1–3). This may influence the results as these agents differ in their efficacy, half-life, mechanism of

bioavailability and elimination/excretion (in stool for example). Infliximab, for instance, is administered intravenously (IV), while adalimumab, certolizumab pegol, and golimumab are administered subcutaneously (SC). Intravenous administration is associated with large volume, rapid central distribution with low variability in bioavailability. Absorption from SC administration is slow, and it may induce more immunogenicity.⁹⁰ The IBDResponse⁶¹ trial has drawn attention to this problem and challenged the results obtained from previous studies where a mix of biological treatment agents were registered.

Time interval between medication and surgical intervention

Most of the studies chose a 8- to 12-week interval from the last dose of a-TNF to the date of surgical surgery (Tables 1–3). This is mainly based on pharmacokinetics of infliximab (the most commonly used drug), assuming first-order elimination

TABLE 4. Meta-analyses and Systematic Reviews That Investigated the Effect of Biological Treatment on Postoperative Outcome in Patients With Inflammatory Bowel Disease

Author/Publication's Year		Disease	Primary Outcome
Systematic/narrative reviews			
1	Subramanian ⁶⁹ et al (2006)	CD and UC	Postoperative complications
2	Ali ⁷⁰ et al (2012)	CD and UC	Postoperative complications
3	El-Hussuna ⁷¹ et al (2014)	CD	Postoperative complications
4	Papaconstantinou ⁷² et al (2014)	CD	Postoperative complications
5	Saab ⁷⁸ et al (2015)	CD	Postoperative complications
6	Holubar ⁷³ et al (2015)	CD and UC	Overall/infectious complications
7	Alexakis ⁷⁴ et al (2015)	UC	Colectomy and hospitalization rates
8	Chang ⁷⁵ et al (2015)	CD	Surgical complications
9	Kotze ⁷⁶ et al (2017)	CD	Postoperative complications
10	Engel ⁷⁷ et al (2017)	CD and UC	Postoperative complications
Meta-analyses			
1	Yang ¹¹⁶ et al (2009)	UC	Postoperative complications
2	Ehteshami-Afshar ⁸² et al (2011)	CD and UC	Colectomy and hospitalization rates
3	Kopylov ⁸¹ et al (2012)	CD	Colectomy and hospitalization rates
4	El-Hussuna ¹⁰ et al (2013)	CD	Anastomotic complications
5	Billioud ⁸⁴ et al (2013)	CD and UC	Postoperative complications
6	Narula ⁸⁵ et al (2013)	CD and UC	Postoperative complications
7	Rosenfeld ⁸⁶ et al (2013)	CD	Postoperative complications
8	Yang ¹¹⁶ et al (2014)	CD	Postoperative complications
9	Ahmed Ali ⁸⁷ et al (2014)	CD	Overall/infectious complications
10	Selvaggi ¹²⁰ et al (2015)	UC	Pouch-related postoperative complications
11	Waterland ⁸⁸ et al (2016)	CD	Infectious complications
12	Law ⁸⁹ et al (2018)	CD and UC	Overall/infectious complications
13	Xu ⁸⁰ et al (2018)	CD	Postoperative complications
14	Yung ⁸¹ et al (2018)	CD and UC	Postoperative complications

It shows 12 reviews about CD, 9 about mixed population, and only 3 about UC.

action, pharmacodynamics and pharmacokinetics including

TABLE 5. Clinical Studies About Risk Factors for Unfavorable Postoperative Outcome in Patients With Crohn Disease and Ulcerative Colitis

Author/Publication's Year	Disease	Cohort/Treated (12,830/2881)	Type of Biological Treatment	Weeks Before Surgery
Observational retrospective studies				
1 Iesalnieks ¹²¹ et al (2008)	CD	282/4	IFX	8
2 Sampietro ¹²² et al (2009)	CD	393/13	Biological agents (not specified)	Not specified
3 Canedo ¹²³ et al (2010)	CD and UC	213/61	IFX, ADA	8
4 Holubar ¹²⁴ et al (2010)	CD	92/32	Biological agents (not specified)	Not specified
5 De Silva ¹²⁵ et al (2011)	UC	666/58	IFX	Not specified
6 Mascarenhas ¹²⁶ et al (2012)	CD	93/19	IFX, ADA and others (not specified)	12
7 Riss ¹²⁷ et al (2012)	CD	182/3	IFX	1
8 Tzivanakis ¹²⁸ et al (2012)	CD	207/not stated	IFX	Not specified
9 Bellolio ¹²⁹ et al (2013)	CD	434/42	Biological agents (not specified)	Not specified
10 Gu ¹³⁰ et al (2013)	UC	204/73	Anti-TNF- α (not specified)	Not specified
11 Bartels ¹³¹ et al (2013)	UC	71/16	Anti-TNF- α (not specified)	Not specified
12 Bewtra ¹³² et al (2013)	UC	830/65	IFX	Not specified
13 Hicks ¹³³ et al (2014)	UC	179/43	IFX	Not specified
14 Morar ¹³⁴ et al (2015)	CD	142/4	IFX, ADA	4
15 Zuo ¹³⁵ et al (2015)	CD	344/8	IFX	Not specified
16 Feuerstein ¹³⁶ et al (2015)	UC	209/24	Anti-TNF- α (not specified)	Not specified
17 Li ¹³⁶ et al (2016)	CD	1461/190	Biological agents (not specified)	Not specified
18 Germain ¹³⁷ et al (2016)	CD	137/13	Anti-TNF- α (not specified)	8
19 Yamamoto ¹³⁸ et al (2016)	CD	231/79	IFX, ADA	8
20 Sahami ¹³⁹ et al (2016)	UC	640/51	Anti-TNF- α (not specified)	12
21 Guo ¹⁴⁰ et al (2017)	CD	118/11	Anti-TNF- α (not specified)	24
22 Collaborative ¹⁰¹ (2017)	CD	375/82	IFX, ADA, CZM, and others (not specified)	12
23 Diederer ¹⁴¹ et al (2017)	UC	422/14	Anti-TNF- α (not specified)	12
24 Galata ¹⁰⁰ et al (2018)	CD	305/72	IFX, ADA, golimumab, vedolizumab, and others	4
25 Heilmann ¹⁴² et al (2018)	CD and UC	1000/71	Biological agents (not specified)	6
Reviews				
26 Huang ¹⁴³ et al (2015)	CD	1833	Biological agents (not specified)	Not specified
27 Beddy ¹⁴⁴ et al (2011)	CD and UC	Not stated	Biological agents (not specified)	Not specified

In these studies, biological treatment was analyzed as part of the risk factors in multivariate analysis.

IFX, infliximab; ADA, adalimumab; CZM, cemiziv; anti-TNF- α , antitumor necrosis factor alpha agents.

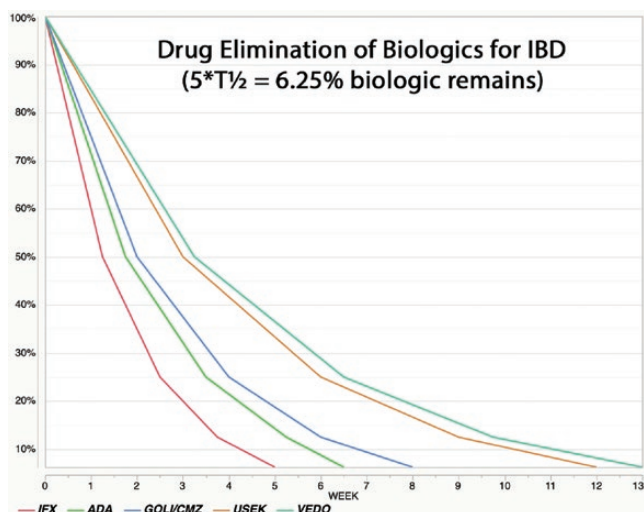


FIGURE 1. Graphical representation of the theoretical in vivo half-lives of biologic agents used to treat CUC. Note this graph assumes first-order elimination pharmacokinetics. With permission from Stefan Holubar.

kinetics (Fig. 1), which may or may not be applicable; of note, trace levels of infliximab can be found up to 12 weeks after administration.⁹⁰ However, during this time, drug concentration and its efficacy can vary from a peak at the time of administration to a trace level at the end of time interval (see the next section).

Drug concentration in the peripheral blood (serum levels of biological agents)

Drug concentration in the peripheral blood correlates with drug concentration in the inflamed tissue⁹¹ and it varies during the 12-week period prior to surgery with a peak at the time of administration and a fading to trace levels in the following weeks (Fig. 1). Drug concentration is essential for drug action and for the stimulation of antidrug antibody formation.⁹⁰ Reporting biologic treatment at 12 weeks interval can therefore be misleading and interpretation of the results may differ. Only 4 studies investigated drug concentration in the peripheral blood.^{29, 30, 60, 61} These 4 studies reported the drug concentration at different time intervals using trough levels in the serum. These trough levels are measured by using enzyme-linked immunosorbent assay, and these levels may be decreased by leakage of the antibodies through inflamed bowel mucosa into the stool⁹² and/or deactivation by development of antibodies against the injected anti-TNF (antidrug antibodies).⁹³ Fumery et al⁶⁰ reported drug concentration of 76 of the 93 patients who received a-TNF treatment within a period of 12 weeks before surgery. The measurement of trough level was done on the day of surgery.⁶⁰ Lau et al reported serum drug concentrations from samples drawn at varying preoperative time points in 143 patients with IBD treated with a-TNF.³⁰

Waterman et al reported drug concentration in a subset of 19 patients with UC exposed to a-TNF treatment within 8 weeks prior to surgery. The authors also reported antidrug antibodies.²⁹ Analysis by Waterman et al included patients exposed to a-TNF treatment within 180 days of surgery. El-Hussuna⁶¹ et al reported in IBDResponse study drug concentration and antidrug antibodies for 18 patients with IBD. Samples were collected within 24 hours before surgery as well as 6, 24, and 48 hours after surgery providing a unique chance to examine the drug and antidrug antibodies in this group of patients.

Antidrug antibodies

The IBDResponse trial⁶¹ showed that some patients develop antidrug antibodies to a-TNF agents regardless of the route of administration (IV or SC). These antidrug antibodies may reduce the efficacy of drugs. Misinterpretation of results in the studies where these antibodies were not measured cannot be dismissed. Only 2 studies^{29, 61} measured and reported these antidrug antibodies. Concurrent administration of immunomodulators has been shown to reduce the formation of antidrug antibodies.⁹⁰

How the Included Studies Adjusted for Potential Confounding Factors in Multivariate Analysis

Many researchers suspected confounding factors play a role and affect postoperative outcomes. These factors have a varying degree of influence on the postoperative outcome and with increasing influence by the number of concurrent risk factors. Some of these factors are well studied such as concurrent medications, whereas others are less studied such as preoperative optimization, the latter which has been recently shown to have a strong influence on the postoperative outcome.^{94, 95} Three categories of confounding factors can be identified and have been studied:

Factors with questionable impact as shown by the studies that adjusted for these factors

- Type of surgical intervention and access to abdominal cavity (studies of ileo-caecal resection,^{15, 19, 34, 60, 96} studies of different types of bowel resections with or without stricturoplasty,^{11–14, 16–18, 22–25, 27–33, 36–40, 42, 59, 66, 97} and studies of ileostomy reversal.²⁶ In patients with UC most of the studies reported postoperative outcome after subtotal colectomy,^{36, 43–46, 48, 55, 67, 68} completion proctectomy,⁵⁷ ileal pouch-anal anastomosis,^{43–45, 48, 50, 51, 53, 54, 58, 98} and other procedures in addition to the above-mentioned.⁴⁹
- One third of studies adjusted for BMI.^{15, 16, 19, 24, 28, 29, 32, 33, 36, 38–40, 43, 45, 49, 53, 54, 58–60, 96}
- Many studies adjusted for American Society of Anesthesiology score.^{14, 15, 33, 34, 36, 38, 45, 54, 59, 96} However, many studies reported comorbidity,^{19, 26–28, 32, 33, 36, 40, 45, 48, 53, 54, 67} but few used Charlson comorbidity score.^{16, 25, 66, 68}

- (d) Previous intestinal resections reported/adjusted for in 14 studies.^{12, 14–16, 19, 23, 27, 28, 36, 39, 40, 60, 96, 99}
- (e) Disease phenotype reported was reported in many studies.^{13–19, 22, 29, 30, 34, 36, 38, 39, 59, 60, 97, 99}
- (f) The affected bowel segment/length or disease location was reported in 20 studies.^{12–15, 19, 27, 29, 36, 39, 44–46, 50, 58–60, 96, 97, 99}
- (g) Duration of operation was reported in some studies.^{15, 23, 28, 36, 40, 42, 53, 59, 96}
- (h) Preoperative intra-abdominal sepsis (intra-abdominal abscess and/or enteric fistula) was reported in 9 studies.^{13, 15, 16, 30, 34, 42, 59, 60, 96}
- (i) Disease duration was reported in 15 studies.^{12, 16, 18, 22, 27, 28, 35, 36, 38, 39, 46, 58, 66, 67, 99}
- (j) Urgency of surgical intervention was reported in one third of studies.^{12, 14–18, 25, 27, 29, 31, 33, 35–38, 40, 42, 45, 48, 50, 54, 59, 60, 68, 96, 97} Some studies excluded urgent/emergency operations to attain a homogenous group of elective surgical procedures.^{28, 38, 61}
- (k) Surgeon's experience (trainee, general surgeon, or colorectal surgeon) was reported in only 3 studies.^{30, 31, 96}
- (l) The type/configuration of anastomosis or stoma construction was reported in several studies.^{14, 15, 19, 33, 45, 50, 54, 59, 96} One study included stricturoplasty in addition to primary anastomosis.¹⁴ Patients who received a diverting stoma were excluded in some studies,^{38, 42} whereas other studies investigated these patients in subgroup analyses.⁵⁹
- (m) Other factors such as intra-operative blood loss,^{37, 42} indication for surgery,^{12, 23, 25, 27, 29, 30, 34, 40, 50} multicenter versus single centre,^{27–30, 32, 33, 35, 39, 42, 54} leucocytosis¹⁰⁰ were reported but they appear to have minimum or no effect on the postoperative outcome. Close cooperation between surgeon and gastroenterologist in presurgical decision making may have an impact on postoperative outcome,^{25, 101} but this factor is difficult to measure and not reported.

Factors expected to have large impact but were less well studied

- (a) Preoperative optimization: few studies reported interventions to optimize the patients prior to surgery, for example, nutritional support,^{15, 59, 60, 96} correction of anemia, or prehabilitation.
- (b) Use of a mechanical bowel preparation, which has been associated with anastomotic leaks rates, was reported in only one study.¹⁵

Factors which would be expected to have a large impact on postoperative outcome

- (a) Concurrent medication was reported by almost all studies^{11–15, 17–19, 22–25, 27–40, 43–46, 48–51, 53–55, 58–60, 66, 67, 96–98} except one population-based study on UC.⁶⁸
- (b) Although nutritional status is widely known to influence postoperative outcomes,^{95, 94} it was only reported in some studies, whereas others did not report it.^{60, 66, 59, 24, 96, 19} Nutritional status was measured indirectly by assessment of serum albumin and/or haemoglobin^{11, 16, 18, 24, 25, 27, 29, 30, 32, 33, 35–37, 39, 40, 45, 49, 50, 55, 59, 60, 96–98} in those studies that adjusted for nutritional status. The IBDResponse trial⁶¹ used a validated standard score (nutritional risk screening

which included weight loss more 10% of body weight) to assess nutritional status.

- (c) Smoking is well-documented risk factor of surgical site infection⁹⁵ however, with small sample size series (as in case of most studies) is it difficult to demonstrate statistical significance due to lack of power. Data on smoking were reported by some studies.^{11, 13–16, 18, 19, 24, 28, 30, 32–36, 46, 49, 50, 55, 58–60, 68, 96, 97}
- (d) Crohn Disease Activity Index (CDAI) was reported in 2 studies only,^{60, 99} whereas Harvey–Bradshaw Index was reported in one study.⁶¹ One study reported ACG severity of disease index,³⁸ whereas another one applied a local classification of disease activity.²⁸ In UC studies, one study used deprivation index,⁶⁸ whereas others used a local disease activity index^{36, 48} or the Mayo score.⁵⁵ Regarding disease severity, there is a high likelihood that the most severely ill patients received anti-TNF, whereas the less ill patients did not. One study tried to compare similar groups where all patients received anti-TNF at any time during the disease course,¹⁴ either at time of surgery or prior to (but withheld) versus after surgery.

How the Included Studies Reported Outcomes

Different methods were used to report the postoperative outcome making the comparison of studies difficult. Some studies reported major and minor complications.^{16, 19, 22, 23, 37, 39} Others reported short-term (early) and long-term (late) postoperative morbidity.^{12, 23, 28} A third category of studies reported septic/infectious^{11, 12, 14, 17, 19, 27, 31, 32, 35, 40, 43, 46, 54, 58, 59, 99} and nonseptic/noninfectious complications. A few studies used the classification of surgical complications versus medical complications,^{18, 30, 44, 97} whereas other studies presented postoperative complications without classification or grading.^{14, 16, 24, 25, 27, 29, 33, 34, 49, 51, 60, 66, 67, 98} However, there was increasing tendency to report outcomes according to Clavien–Dindo classification of postoperative complications.^{14, 19, 28, 30, 53, 59, 60, 96, 97} One study reported outcome when biological treatment was used after surgical intervention in CD.⁹⁹

To the best of our knowledge, one study has used the Comprehensive Complications Index, a relatively new composite outcome of any complication weighted by Clavien–Dindo level.¹²⁰ Interestingly, no difference in the length of stay (LOS) postoperatively was shown between patients treated with biological agents and those who did not receive treatment in the studies where LOS was reported. This was unexpected in studies that reported increased complication rates in patients treated with biological agents, as LOS will be longer in patients with postoperative complications.¹⁴⁵

How the Different Studies Conducted Statistical Analyses

Most of the statistical analyses were done in similar fashion, that is, univariate analyses with chi-square test, Student *t* test, or Fisher exact test for categorical and Mann–Whitney,

Wilcoxon signed-rank tests for continuous variables. However, regarding multivariable analyses, many studies did not report what variables were entered in multivariate analyses, nor how these variables were chosen for the multivariate analyses.^{23, 38} Errors in interpretation of statistical results were not uncommon for instance lack of adjustment for confounding factors.^{26,37}

DISCUSSION

Despite the large number of studies available in the literature, the relationship between biologic treatment and postoperative outcome in IBD is still controversial. During the last 15 years, there have been improvements in study design, statistical analyses, and sample size, moving from high risk of bias studies³⁷ to more recently low risk of bias studies⁵⁹; nevertheless, the topical debate and controversy continues. There are 3 layers of complexity that made it difficult to reach definitive conclusions about the issue at hand:

1. Difficulty of conducting clinical research compared with basic science research. Clinical research is becoming more difficult due to the increased complexity of regulations and governance which can be far from patients' interests.¹⁰² Up to half of the approved studies by ethics committees are never published.¹⁰³ It is well documented that clinical research attracts much less funding than basic science or translational research adding another challenge for outcomes research.¹⁰⁴
2. Difficulty in conducting research in surgery compared with medical specialties. Variation in surgical practice affects postoperative outcome and leads to a strong confounding factor in surgical research. Few studies include surgeon experience or years from training.
3. Difficulty in conducting IBD research. There is no doubt IBD is complex and heterogeneous; the treatment is complex as are the preoperative and postoperative assessments. To this end, a number of groups have been working toward developed standardized outcome sets to facilitate comparison of data and effective meta-analysis.¹⁰⁵⁻¹⁰⁸ Moreover, investigator-initiated trials often fail due to insufficient enrollment of patients with IBD,¹⁰⁰ and a priori power analyses are rarely reported. Only one third to half of patients with IBD need surgery during their lifetime (75% in CD⁸) making it even harder to recruit patients. Funding is a general problem in research, but it is more prominent in investigator-initiated trials, especially in IBD. Having said this, it might be assumed that research in IBD surgery is very well planned to reduce poor quality and ensure best use of resources. However, as of today, this is not the case which as this review demonstrates. Large amount of resources were used in repeating studies with minor variations in design.

Crohn Disease

According to recently published guidelines,¹⁰⁹⁻¹¹¹ surgery in patients with receiving a-TNF therapy may be associated with an increased risk of complications. Chronologically, the ASCRS 2015 clinical practice guideline on CD stated that

patient receiving preoperative biologic treatment (ie, a-TNF or cyclosporine) should be considered for staged procedures because of postoperative complications risk.¹⁰⁹ The authors suggested final decision should be up to surgeon discretion using an individualized approach to each patient. A delay of at least 8 weeks was proposed for elective whenever possible. The recommendations were graded as weak, quality of evidence 2C.

The ECCO published its third evidence-based CD consensus in 2016,¹¹⁰ where the impact of biologic treatment for patients undergoing surgery was deemed unclear and controversial. The authors based their statement on controversial data and advocated optimal preoperative preparation. A joint statement of the ESCP and ECCO considered a-TNF to be associated with risk of postoperative complications, particularly sepsis (surgical site infections, abdominal abscesses, anastomotic leaks) and higher readmissions.¹¹¹ Nevertheless, no recommendation was made regarding the interval of biological treatment withdrawal. All guidelines agree on the higher risk associated with long-term steroid therapy. Prednisolone 20 mg daily or higher for >6 weeks was associated with higher postoperative surgical complications, especially when used in combination with biologic treatment.¹⁰⁹⁻¹¹¹ The American Gastroenterological Association (AGA) did not published specific recommendation on the clinical management of biological treatment prior to surgery.

Ulcerative Colitis

Published guidelines on the surgical management of UC includes the ECCO consensus.¹¹² For acute situations, performing staged procedures (ie, subtotal colectomy with end ileostomy as first stage)¹¹⁹ was advised for patients receiving biologic treatment (a-TNF) and/or Prednisolone 20 mg daily or higher for >6 weeks. Regarding preoperative management of biologic treatments, the increased risks of postoperative complications, although controversial, were outlined, and single-stage procto-colectomy with ileo-anal pouch reconstruction, for patients under a-TNF therapy, was not recommended.¹¹² Guidelines from the ASCRS published in 2014¹¹³ lacked definitive consensus statement on the management of a-TNF prior to elective surgery for UC. According to the authors, literature was insufficient to assess the impact of biological treatment on postoperative outcomes and there was a claim for multi-institutional larger studies.

There is no statistical model to predict the effect of various confounding factors on the postoperative outcomes in patients with IBD. It is nevertheless believed that the weight of these confounding factors in the final model is certainly different, and no standard recommendations for which variables to include in a multivariate model exists; thus, there is wide heterogeneity in model building, thus leading to varying and potentially noncomparable results and conclusions. Going forward, propensity-score matching or inclusion of the propensity

score in the multivariate model, and other novel methods such as the difference-in-difference and use of instrumental variables all may play a role in reducing both the measured, and unmeasured, bias and confounding in these studies.

Future Perspectives

ESCP conducted a snapshot audit in 2015 in which patients with CD undergoing ileo-caecal resection and right-side colectomy were included, but this snapshot study did not provide a definitive answer regarding the effect of biologic treatment on postoperative outcomes.¹⁰¹ Clearly, there is a need for randomized controlled trials investigating the effect of biological treatment on postoperative outcome in patients with IBD. The Pre-operative Continuation versus Discontinuation of anti-TNF treatment in Patients with Crohn's Disease (PCDantiPCD) trial protocol was presented in the 16th Nordic postgraduate course in colorectal surgery and at the ESCP 2018 trial session, and integrates many of the points discussed above.

Measurement of drug concentration, antidrug antibodies, and application of a standardized validated scoring systems for disease activity, nutritional status, and smoking will lead to better understanding of the effect/weight of different covariates in a model that describes how anti-TNF treatment influence the postoperative outcome. Meta-analysis of these randomized controlled trial will provide a solid evidence to eliminate the uncertainty of previous observational studies.

Limitations

This narrative updated review has the limitations of the studies included. It was not planned as a new systematic review or meta-analysis; therefore, no statistical analysis was performed.

CONCLUSION

Many studies have investigated the effect of biologic treatment on postoperative outcome using different methodological approaches (retrospective, prospective, population-based, experimental, snapshot audit, and meta-analyses) with divergent results. Future studies should focus on the avoiding the above highlighted weakness of the studies we reviewed. Consensus guidelines by the invested societies, such as ECCO, Crohn's & Colitis Foundation (formerly known as the Crohn's & Colitis Foundation of America [CCFA]), and ESCP, are needed to guide future research. There is also a need for a randomized controlled trial to define the association, or lack thereof, between biological and adverse postoperative outcomes.

SUPPLEMENTARY DATA

Supplementary data are available at *Crohn's & Colitis 360* online.

REFERENCES

- Behm B, Babilas P, Landthaler M, et al. Cytokines, chemokines and growth factors in wound healing. *J Eur Acad Dermatol Venereol*. 2012;26:812–820.
- Lin E, Lowry SF. Inflammatory cytokines in major surgery: a functional perspective. *Intensive Care Med*. 1999;25:255–257.
- Park JE, Barbul A. Understanding the role of immune regulation in wound healing. *Am J Surg*. 2004;187:115–165.
- Tsirogianni AK, Moutsopoulos NM, Moutsopoulos HM. Wound healing: immunological aspects. *Injury*. 2006;37 (Suppl 1):S5–S12.
- Peyrin-Biroulet L, Harmsen WS, Tremaine WJ, Zinsmeister AR, Sandborn WJ, Loftus EV Jr. Surgery in a population-based cohort of Crohn's disease from Olmsted county 1970–2004. *Am J Gastroenterol*. 2012;13:1133–1145.
- Burisch J, Kiudelis G, Kupcinskas L, et al. Natural disease course of Crohn's disease during the first 5 years after diagnosis in a European population-based inception cohort: an Epi-IBD study. *Gut*. 2018. pii: gutjnl-2017-315568. doi:10.1136/gutjnl-2017-315568. [Epub ahead of print].
- Frolkis AD, Lipton DS, Fiess KM, et al. Cumulative incidence of second intestinal resection in Crohn's disease: a systematic review and meta-analysis of population-based studies. *Am J Gastroenterol*. 2014;109:1739–1748.
- Peyrin-Biroulet L, Oussalah A, Williet N, et al. Impact of azathioprine and tumour necrosis factor antagonists on the need for surgery in newly diagnosed Crohn's disease. *Gut*. 2011;60:930–936.
- Colombel JF, Ricart E, Loftus EV Jr, et al. Management of Crohn's disease of the ileoanal pouch with infliximab. *Am J Gastroenterol*. 2003;98:2239–2244.
- El-Hussuna A, Krag A, Olaison G, et al. The effect of anti-tumour necrosis factor alpha agents on postoperative anastomotic complications in Crohn's disease: a systematic review. *Dis Colon Rectum*. 2013;56:1423–1433.
- Tay GS, Binion DG, Eastwood D, et al. Multivariate analysis suggests improved perioperative outcome in Crohn's disease patients receiving immunomodulator therapy after segmental resection and/or strictureplasty. *Surgery*. 2003;134:565–572; discussion 572.
- Colombel JF, Loftus EV Jr, Tremaine WJ, et al. Early postoperative complications are not increased in patients with Crohn's disease treated perioperatively with infliximab or immunosuppressive therapy. *Am J Gastroenterol*. 2004;99:878–883.
- White EC, Melmed GY, Vasiliauskas E, et al. Does preoperative immunosuppression influence unplanned hospital readmission after surgery in patients with Crohn's disease? *Dis Colon Rectum*. 2012;55:563–568.
- Myrelid P, Marti-Gallostra M, Ashraf S, et al. Complications in surgery for Crohn's disease after preoperative antitumour necrosis factor therapy. *Br J Surg*. 2014;101:539–545.
- Serradori T, Germain A, Scherrer ML, et al. The effect of immune therapy on surgical site infection following Crohn's disease resection. *Br J Surg*. 2013;100:1089–1093.
- Syed A, Cross RK, Flasar MH. Anti-tumour necrosis factor therapy is associated with infections after abdominal surgery in Crohn's disease patients. *Am J Gastroenterol*. 2013;108:583–593.
- Bafford AC, Powers S, Ha C, et al. Immunosuppressive therapy does not increase operative morbidity in patients with Crohn's disease. *J Clin Gastroenterol*. 2013;47:491–495.
- Kotze PG, Saab MP, Saab B, et al. Tumour necrosis factor alpha inhibitors did not influence postoperative morbidity after elective surgical resections in Crohn's disease. *Dig Dis Sci*. 2017;62:456–464.
- Jouvin I, Lefevre JH, Creavin B, et al. Saint-Antoine IBD Network. Postoperative morbidity risks following ileocolic resection for Crohn's disease treated with anti-TNF alpha therapy: a retrospective study of 360 patients. *Inflamm Bowel Dis*. 2018;24:422–432.
- Kotze PG, Ludvig JC, Teixeira FV, et al. Disease duration did not influence the rates of loss of efficacy of the anti-TNF therapy in Latin American Crohn's disease patients. *Digestion*. 2015;91:158–163.
- Shim HH, Ma C, Kotze PG, et al. Preoperative ustekinumab treatment is not associated with increased postoperative complications in Crohn's disease: a Canadian Multi-Centre Observational Cohort Study. *J Can Assoc Gastroenterol*. 2018;1:115–123.
- Zimmerman LA, Saites CG, Bairdain S, et al. Postoperative complications in children with Crohn disease treated with infliximab. *J Pediatr Gastroenterol Nutr*. 2016;63:352–356.
- Marchal L, D'Haens G, Van Assche G, et al. The risk of post-operative complications associated with infliximab therapy for Crohn's disease: a controlled cohort study. *Aliment Pharmacol Ther*. 2004;19:749–754.
- Lightner AL, McKenna NP, Tse CS, et al. Postoperative outcomes in ustekinumab-treated patients undergoing abdominal operations for Crohn's disease. *J Crohns Colitis*. 2018;12:402–407.
- Kunitake H, Hodin R, Shellito PC, et al. 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; discussion 1736.
- Regadas FS, Pinto RA, Murad-Regadas SM, et al. Short-term outcome of infliximab and other medications on patients with inflammatory bowel disease undergoing ileostomy reversal. *Colorectal Dis*. 2011;13:555–560.

27. Rizzo G, Armuzzi A, Pugliese D, et al. Anti-TNF- α therapies do not increase early postoperative complications in patients with inflammatory bowel disease. An Italian single-center experience. *Int J Colorectal Dis.* 2011;26:1435–1444.
28. Krane MK, Allaix ME, Zoccali M, et al. Preoperative infliximab therapy does not increase morbidity and mortality after laparoscopic resection for inflammatory bowel disease. *Dis Colon Rectum.* 2013;56:449–457.
29. Waterman M, Xu W, Dinani A, et al. Preoperative biological therapy and short-term outcomes of abdominal surgery in patients with inflammatory bowel disease. *Gut.* 2013;62:387–394.
30. Lau C, Dubinsky M, Melmed G, et al. The impact of preoperative serum anti-TNF α therapy levels on early postoperative outcomes in inflammatory bowel disease surgery. *Ann Surg.* 2015;261:487–496.
31. Alsaleh K, Gaidos JK, Kang L, et al. Timing of last preoperative dose of infliximab does not increase postoperative complications in inflammatory bowel disease patients. *Dig Dis Sci.* 2016;61:2602–2607.
32. Lightner AL, Raffals LE, Mathis KL, et al. Postoperative outcomes in vedolizumab-treated patients undergoing abdominal operations for inflammatory bowel disease. *J Crohns Colitis.* 2017;11:185–190.
33. Shwaartz K, Fields AC, Sobrero M, et al. Effect of anti-TNF agents on postoperative outcomes in inflammatory bowel disease patients: a single institution experience. *J Gastrointest Surg.* 2016;20:1636–1642.
34. Appau KA, Fazio VW, Shen B, et al. Use of infliximab within 3 months of ileocolonic resection is associated with adverse postoperative outcomes in Crohn's patients. *J Gastrointest Surg.* 2008;12:1738–1744.
35. Yamada A, Komaki Y, Patel N, et al. Risk of postoperative complications among inflammatory bowel disease patients treated preoperatively with vedolizumab. *Am J Gastroenterol.* 2017;112:1423–1429.
36. Uchino M, Ikeuchi H, Matsuoka H, et al. Risk factors for surgical site infection and association with infliximab administration during surgery for Crohn's disease. *Dis Colon Rectum.* 2013;56:1156–1165.
37. Indar AA, Young-Fadok TM, Heppell J, et al. Effect of perioperative immunosuppressive medication on early outcome in Crohn's disease patients. *World J Surg.* 2009;33:1049–1052.
38. Nasir BS, Dozois EJ, Cima RR, et al. Perioperative anti-tumor necrosis factor therapy does not increase the rate of early postoperative complications in Crohn's disease. *J Gastrointest Surg.* 2010;14:1859–1865; discussion 1865.
39. Kasperek MS, Bruckmeier A, Beigel F, et al. Infliximab does not affect postoperative complication rates in Crohn's patients undergoing abdominal surgery. *Inflamm Bowel Dis.* 2012;18:1207–1213.
40. Canedo J, Lee SH, Pinto R, et al. Surgical resection in Crohn's disease: is immunosuppressive medication associated with higher postoperative infection rates? *Colorectal Dis.* 2011;13:1294–1298.
41. Regueiro M, El-Hachem S, Kip KE, et al. Postoperative infliximab is not associated with an increase in adverse events in Crohn's disease. *Dig Dis Sci.* 2011;56:3610–3615.
42. El-Hussuna A, Andersen J, Bisgaard T, et al. Biologic treatment or immunomodulation is not associated with postoperative anastomotic complications in abdominal surgery for Crohn's disease. *Scand J Gastroenterol.* 2012;47:662–668.
43. 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;204:956–962; discussion 962.
44. 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.
45. Gu J, Remzi FH, Shen B, et al. Operative strategy modifies risk of pouch-related outcomes in patients with ulcerative colitis on preoperative anti-tumor necrosis factor- α therapy. *Dis Colon Rectum.* 2013;56:1243–1252.
46. Nelson R, Liao C, Fichera A, et al. Rescue therapy with cyclosporine or infliximab is not associated with an increased risk for postoperative complications in patients hospitalized for severe steroid-refractory ulcerative colitis. *Inflamm Bowel Dis.* 2014;20:14–20.
47. Zittan E, Milgrom R, Ma GW, et al. Preoperative anti-tumor necrosis factor therapy in patients with ulcerative colitis is not associated with an increased risk of infectious and noninfectious complications after ileal pouch-anal anastomosis. *Inflamm Bowel Dis.* 2016;22:2442–2447.
48. Kulaylat AS, Kulaylat AN, Schaefer EW, et al. Association of preoperative anti-tumor necrosis factor therapy with adverse postoperative outcomes in patients undergoing abdominal surgery for ulcerative colitis. *JAMA Surg.* 2017;152:e171538.
49. Lightner AL, McKenna NP, Moncrief S, et al. Surgical outcomes in vedolizumab-treated patients with ulcerative colitis. *Inflamm Bowel Dis.* 2017;23:2197–2201.
50. Ferrante M, de Buck van Overstraeten A, Schils N, et al. Perioperative use of vedolizumab is not associated with postoperative infectious complications in patients with ulcerative colitis undergoing colectomy. *J Crohns Colitis.* 2017;11:1353–1361.
51. Mor IJ, Vogel JD, da Luz Moreira A, et al. Infliximab in ulcerative colitis is associated with an increased risk of postoperative complications after restorative proctocolectomy. *Dis Colon Rectum.* 2008;51:1202–1207; discussion 1207.
52. Ferrante M, D'Hoore A, Vermeire S, et al. Corticosteroids but not infliximab increase short-term postoperative infectious complications in patients with ulcerative colitis. *Inflamm Bowel Dis.* 2009;15:1062–1070.
53. 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.
54. Gainsbury ML, Chu DI, Howard LA, et al. Preoperative infliximab is not associated with an increased risk of short-term postoperative complications after restorative proctocolectomy and ileal pouch-anal anastomosis. *J Gastrointest Surg.* 2011;15:397–403.
55. Bregnbak D, Mortensen C, Bendtsen F. Infliximab and complications after colectomy in patients with ulcerative colitis. *J Crohns Colitis.* 2012;6:281–286.
56. Jones GR, Kennedy NA, Lees CW, et al. Systematic review: the use of thiopurines or anti-TNF in post-operative Crohn's disease maintenance – progress and prospects. *Aliment Pharmacol Ther.* 2014;39:1253–1265.
57. Uchino M, Ikeuchi H, Matsuoka H, et al. Infliximab administration prior to surgery does not increase surgical site infections in patients with ulcerative colitis. *Int J Colorectal Dis.* 2013;28:1295–1306.
58. Eshuis EJ, Al Saady RL, Stokkers PC, et al. Previous infliximab therapy and postoperative complications after proctocolectomy with ileum pouch anal anastomosis. *J Crohns Colitis.* 2013;7:142–149.
59. Brouquet A, Maggiori L, Zerbib P, et al. GETAID Chirurgie Group. Anti-TNF therapy is associated with an increased risk of postoperative morbidity after surgery for ileocolonic Crohn disease: results of a prospective nationwide cohort. *Ann Surg.* 2018;267:221–228.
60. Fumery M, Seksik P, Auzolle C, et al.; REMIND Study Group Investigators. Postoperative complications after ileocecal resection in Crohn's disease: a prospective study from the REMIND Group. *Am J Gastroenterol.* 2017;112:337–345.
61. El-Hussuna A, Qvist N, Zangenberg MS, et al. No effect of anti-TNF- α agents on the surgical stress response in patients with inflammatory bowel disease undergoing bowel resections: a prospective multi-center pilot study. *BMC Surg.* 2018;18:91.
62. Ploug T, Andersen K, Hansen K, et al. Influence of adalimumab treatment on anastomotic strength, degree of inflammation, and collagen formation: an experimental study on the small intestine of rabbits. *Inflamm Bowel Dis.* 2013;19:254–258.
63. Strebel K, Nielsen SR, Biagini M, et al. Effect of Humira® on intestinal anastomotic response in rabbits. *J Invest Surg.* 2015;28:167–172.
64. Myreliid P, Salim SY, Darby T, et al. Effects of anti-inflammatory therapy on bursting pressure of colonic anastomosis in murine dextran sulfate sodium induced colitis. *Scand J Gastroenterol.* 2015;50:991–1001.
65. Ågren MS, Andersen TL, Andersen L, et al. Nonselective matrix metalloproteinase but not tumor necrosis factor- α inhibition effectively preserves the early critical colon anastomotic integrity. *Int J Colorectal Dis.* 2011;26:329–337.
66. Nørgård BM, Nielsen J, Qvist N, et al. Pre-operative use of anti-TNF- α agents and the risk of post-operative complications in patients with Crohn's disease – a nationwide cohort study. *Aliment Pharmacol Ther.* 2013;37:214–224.
67. Nørgård BM, Nielsen J, Qvist N, et al. Pre-operative use of anti-TNF- α agents and the risk of post-operative complications in patients with ulcerative colitis – a nationwide cohort study. *Aliment Pharmacol Ther.* 2012;35:1301–1309.
68. Ward ST, Mytton J, Henderson L, et al. Anti-TNF therapy is not associated with an increased risk of post-colectomy complications, a population-based study. *Colorectal Dis.* 2018;20:416–423.
69. Subramanian V, Pollok RC, Kang JY, et al. Systematic review of postoperative complications in patients with inflammatory bowel disease treated with immunomodulators. *Br J Surg.* 2006;93:793–799.
70. Ali T, Yun L, Rubin DT. Risk of post-operative complications associated with anti-TNF therapy in inflammatory bowel disease. *World J Gastroenterol.* 2012;18:197–204.
71. El-Hussuna A, Theede K, Olaison G. Increased risk of post-operative complications in patients with Crohn's disease treated with anti-tumour necrosis factor α agents – a systematic review. *Dan Med J.* 2014;61:A4975.
72. Papaconstantinou I, Zeglinas C, Gazouli M, et al. The impact of peri-operative anti-TNF treatment on anastomosis-related complications in Crohn's disease patients. A critical review. *J Gastrointest Surg.* 2014;18:1216–1224.
73. Holubar SD, Holder-Murray J, Flasar M, et al. Anti-tumor necrosis factor- α antibody therapy management before and after intestinal surgery for inflammatory bowel disease: a CCA position paper. *Inflamm Bowel Dis.* 2015;21:2658–2672.
74. Alexakis C, Pollok RC. Impact of thiopurines and anti-tumour necrosis factor therapy on hospitalisation and long-term surgical outcomes in ulcerative colitis. *World J Gastrointest Surg.* 2015;7:360–369.

75. Chang MI, Cohen BL, Greenstein AJ. A review of the impact of biologics on surgical complications in Crohn's disease. *Inflamm Bowel Dis*. 2015;21:1472–1477.
76. Kotze PG, Ghosh S, Bemelman WA, et al. Preoperative use of anti-tumor necrosis factor therapy in Crohn's disease: promises and pitfalls. *Intest Res*. 2017;15:160–165.
77. Engel T, Ungar B, Yung DE, et al. Vedolizumab in IBD—lessons from real-world experience; a systematic review and pooled analysis. *J Crohns Colitis*. 2018;12:245–257.
78. Saab M, Saab B, Olandoski M, et al. Impacto dos anti-TNF nas complicações pós-operatórias na doença de Crohn: Uma revisão. *J Coloproctol*. 2015;35:128–136.
79. Yang ZP, Hong L, Wu Q, et al. Preoperative infliximab use and postoperative complications in Crohn's disease: a systematic review and meta-analysis. *Int J Surg*. 2014;12:224–230.
80. Xu Y, Yang L, An P, Zhou B, Liu G. Meta-analysis: The influence of preoperative infliximab use on postoperative complications of Crohn's disease. *Inflamm Bowel Dis*. 2019;25:261–269. doi:10.1093/ibd/izy246.
81. Yung DE, Horesh N, Lightner AL, et al. Systematic review and meta-analysis: vedolizumab and postoperative complications in inflammatory bowel disease. *Inflamm Bowel Dis*. 2018;24:2327–2338.
82. Ehteshami-Afshar S, Nikfar S, Rezaie A, et al. A systematic review and meta-analysis of the effects of infliximab on the rate of colectomy and post-operative complications in patients with inflammatory bowel disease. *Arch Med Sci*. 2011;7:1000–1012.
83. Kopylov U, Ben-Horin S, Zmora O, et al. Anti-tumor necrosis factor and post-operative complications in Crohn's disease: systematic review and meta-analysis. *Inflamm Bowel Dis*. 2012;18:2404–2413.
84. Billioud V, Ford AC, Tedesco ED, et al. Preoperative use of anti-TNF therapy and postoperative complications in inflammatory bowel diseases: a meta-analysis. *J Crohns Colitis*. 2013;7:853–867.
85. Narula N, Charleton D, Marshall JK. Meta-analysis: peri-operative anti-TNF α treatment and post-operative complications in patients with inflammatory bowel disease. *Aliment Pharmacol Ther*. 2013;37:1057–1064.
86. Rosenfeld G, Qian H, Bressler B. The risks of post-operative complications following pre-operative infliximab therapy for Crohn's disease in patients undergoing abdominal surgery: a systematic review and meta-analysis. *J Crohns Colitis*. 2013;7:868–877.
87. Ahmed Ali U, Martin ST, Rao AD, et al. Impact of preoperative immunosuppressive agents on postoperative outcomes in Crohn's disease. *Dis Colon Rectum*. 2014;57:663–674.
88. Waterland P, Athanasiou T, Patel H. Post-operative abdominal complications in Crohn's disease in the biological era: systematic review and meta-analysis. *World J Gastrointest Surg*. 2016;8:274–283.
89. Law CCY, Narula A, Lightner AL, et al. Systematic review and meta-analysis: preoperative vedolizumab treatment and postoperative complications in patients with inflammatory bowel disease. *J Crohns Colitis*. 2018;12:538–545.
90. Vande Casteele N, Gils A. Pharmacokinetics of anti-TNF monoclonal antibodies in inflammatory bowel disease: adding value to current practice. *J Clin Pharmacol*. 2015;55 Suppl 3:S39–50. doi:10.1002/jcph.374.
91. Yoshihara T, Shinzaki S, Kawai S, et al. Tissue drug concentrations of anti-tumor necrosis factor agents are associated with the long-term outcome of patients with Crohn's disease. *Inflamm Bowel Dis*. 2017;23:2172–2179.
92. Brandse JF, Van Den Brink GR, Wildenberg ME, et al. Loss of infliximab into feces is associated with lack of response to therapy in patients with severe ulcerative colitis. *Gastroenterology*. 2015;149:350–355.e2.
93. Gorovits B, Baltrukonis DJ, Bhattacharya I, et al. Immunoassay methods used in clinical studies for the detection of anti-drug antibodies to adalimumab and infliximab. *Clin Exp Immunol*. 2018;192:348–365.
94. El-Hussuna A, Iesalnieks I, Horesh N, et al. The effect of pre-operative optimization on post-operative outcome in Crohn's disease resections. *Int J Colorectal Dis*. 2017;32:49–56.
95. Zangenberg MS, Horesh N, Kopylov U, et al. Preoperative optimization of patients with inflammatory bowel disease undergoing gastrointestinal surgery: a systematic review. *Int J Colorectal Dis*. 2017;32:1663–1676.
96. 2015 European Society of Coloproctology collaborating group. Risk factors for unfavourable postoperative outcome in patients with Crohn's disease undergoing right hemicolectomy or ileocaecal resection. An international audit by ESCP and S-ECCO. *Colorectal Dis*. 2017. doi:10.1111/codi.13889. [Epub ahead of print].
97. Kotze PG, Magro DO, Martinez CAR, et al. Adalimumab and postoperative complications of elective intestinal resections in Crohn's disease: a propensity score case-matched study. *Color Dis*. 2018;20:211–218.
98. Kennedy R, Potter DD, Moir C, et al. Pediatric chronic ulcerative colitis: does infliximab increase post-ileal pouch anal anastomosis complications? *J Pediatr Surg*. 2012;47:199–203.
99. Regueiro M, El-Hachem S, Kip KE, et al. Postoperative infliximab is not associated with an increase in adverse events in Crohn's disease. *Dig Dis Sci*. 2011;56:3610–3615.
100. Galata C, Weiss C, Hardt J, et al. Risk factors for early postoperative complications and length of hospital stay in ileocecal resection and right hemicolectomy for Crohn's disease: a single-center experience. *Int J Colorectal Dis*. 2018;33:937–945.
101. 2015 European Society of Coloproctology Collaborating Group. Risk factors for unfavourable postoperative outcome in patients with Crohn's disease undergoing right hemicolectomy or ileocaecal resection. An international audit by ESCP and S-ECCO. *Color Dis*. 2018;20:219–227.
102. Salman RA, Beller E, Kagan J, et al. NIH public access regulation and management. *Lancet*. 2014;383:176–185.
103. Blümle A, Meerpohl JJ, Schumacher M, et al. Fate of clinical research studies after ethical approval – follow-up of study protocols until publication. *PLoS One*. 2014;9:e87184.
104. Rygård SL, Kjær MN, Perner A. Statens investering i kliniske forsøg. *Ugeskr Læger*. 2018;180:V09170645.
105. Kim AH, Roberts C, Feagan BG, et al. Developing a standard set of patient-centred outcomes for inflammatory bowel disease – an international, cross-disciplinary consensus. *J Crohns Colitis*. 2018;12:408–418.
106. Ma C, Panaccione R, Fedorak RN, et al. A systematic review for the development of a core outcome set for ulcerative colitis clinical trials. *Clin Gastroenterol Hepatol*. 2017;16:637–647.e13.
107. Schlessinger DI, Iyengar S, Yanes AF, et al. Development of a core outcome set for clinical trials in basal cell carcinoma: study protocol for a systematic review of the literature and identification of a core outcome set using a Delphi survey. *Trials*. 2017;18:490.
108. Sahnan K, Tozer PJ, Adegbola SO, et al.; ENiGMA Collaborators. Developing a core outcome set for fistulising perianal Crohn's disease. *Gut*. 2019;68:226–238.
109. Strong S, Steele SR, Boutsos M, et al.; Clinical Practice Guidelines Committee of the American Society of Colon and Rectal Surgeons. Clinical practice guideline for the surgical management of Crohn's disease. *Dis Colon Rectum*. 2015;58:1021–1036.
110. Gionchetti P, Dignass A, Danese S, et al.; ECCO. 3rd European evidence-based consensus on the diagnosis and management of Crohn's disease 2016: part 2: surgical management and special situations. *J Crohns Colitis*. 2017;11:135–149.
111. Bemelman WA, Warusavitarne J, Sampietro GM, et al. ECCO-ESCP consensus on surgery for Crohn's disease. *J Crohns Colitis*. 2018;12:1–16.
112. Magro F, Gionchetti P, Eliakim R, et al.; European Crohn's and Colitis Organisation (ECCO). Third European evidence-based consensus on diagnosis and management of ulcerative colitis. Part 1: definitions, diagnosis, extra-intestinal manifestations, pregnancy, cancer surveillance, surgery, and ileo-anal pouch disorders. *J Crohns Colitis*. 2017;11:649–670.
113. Ross H, Steele SR, Varma M, et al.; Standards Practice Task Force of the American Society of Colon and Rectal Surgeons. Practice parameters for the surgical treatment of ulcerative colitis. *Dis Colon Rectum*. 2014;57:5–22.
114. Claxton K, Griffin S, Koffijberg H, et al. How to estimate the health benefits of additional research and changing clinical practice. *BMJ*. 2015;351:h5987.
115. Yang Z, Wu Q, Wu K, et al. Meta-analysis: pre-operative infliximab treatment and short-term post-operative complications in patients with ulcerative colitis. *Aliment Pharmacol Ther*. 2010;31:486–492.
116. Yang ZP, Hong L, Wu Q, et al. Preoperative infliximab use and postoperative complications in Crohn's disease: a systematic review and meta-analysis. *Int J Surg*. 2014;12:224–230.
117. Hajibandeh S, Hajibandeh S, Kennedy-Dalby A, et al. Purse-string skin closure versus linear skin closure techniques in stoma closure: a comprehensive meta-analysis with trial sequential analysis of randomised trials. *Int J Colorectal Dis*. 2018;33:1319–1332.
118. Clavien PA, Pugh MA. Measuring and achieving the best possible outcomes in surgery. *Br J Surg*. 2017;104:1121–1122.
119. Yang Z, Wu Q, Wu K, Fan D. Meta-analysis: pre-operative infliximab treatment and short-term post-operative complications in patients with ulcerative colitis. *Aliment Pharmacol Ther*. 2010;31:486–492.
120. Selvaggi F, Pellino G, Canonico S, et al. Effect of preoperative biologic drugs on complications and function after restorative proctocolectomy with primary ileal pouch formation: systematic review and meta-analysis. *Inflamm Bowel Dis*. 2015;21:79–92.
121. Iesalnieks I, Kilger A, Glass H, et al. Intraabdominal septic complications following bowel resection for Crohn's disease: detrimental influence on long-term outcome. *Int J Colorectal Dis*. 2008;23:1167–1174.
122. Sampietro GM, Corsi F, Maconi G, et al. Prospective study of long-term results and prognostic factors after conservative surgery for small bowel Crohn's disease. *Clin Gastroenterol Hepatol*. 2009;7:183–191; quiz 125.
123. Canedo J, Pinto RA, Regadas S, et al. Laparoscopic surgery for inflammatory bowel disease: does weight matter? *Surg Endosc*. 2010;24:1274–1279.
124. Holubar SD, Dozois EJ, Privitera A, et al. Minimally invasive colectomy for Crohn's colitis: a single institution experience. *Inflamm Bowel Dis*. 2010;16:1940–1946.
125. de Silva S, Ma C, Proulx MC, et al. Postoperative complications and mortality following colectomy for ulcerative colitis. *Clin Gastroenterol Hepatol*. 2011;9:972–980.

126. Mascarenhas C, Nunoo R, Asgeirsson T, et al. Outcomes of ileocolic resection and right hemicolectomies for Crohn's patients in comparison with non-Crohn's patients and the impact of perioperative immunosuppressive therapy with biologics and steroids on inpatient complications. *Am J Surg*. 2012;203:375–378; discussion 378.
127. Riss S, Bittermann C, Schwameis K, et al. Determinants for postoperative complications after laparoscopic intestinal resection for Crohn's disease. *Surg Endosc*. 2012;26:933–938.
128. Tzivanakis A, Singh JC, Guy RJ, et al. Influence of risk factors on the safety of ileocolic anastomosis in Crohn's disease surgery. *Dis Colon Rectum*. 2012;55:558–562.
129. Bellolio F, Cohen Z, Macrae HM, et al. Outcomes following surgery for perforating Crohn's disease. *Br J Surg*. 2013;100:1344–1348.
130. Gu J, Stocchi L, Remzi F, et al. Factors associated with postoperative morbidity, reoperation and readmission rates after laparoscopic total abdominal colectomy for ulcerative colitis. *Colorectal Dis*. 2013;15:1123–1129.
131. Bartels SA, Gardenbroek TJ, Bos L, et al. Prolonged preoperative hospital stay is a risk factor for complications after emergency colectomy for severe colitis. *Colorectal Dis*. 2013;15:1392–1398.
132. Bewtra M, Newcomb CW, Wu Q, et al. Mortality associated with medical therapy versus elective colectomy in ulcerative colitis: a cohort study. *Ann Intern Med*. 2015;163:262–270.
133. Hicks CW, Hodin RA, Bordeianou L. Semi-urgent surgery in hospitalized patients with severe ulcerative colitis does not increase overall J-pouch complications. *Am J Surg*. 2014;207:281–287.
134. Morar PS, Hodgkinson JD, Thalayasingam S, et al. Determining predictors for intra-abdominal septic complications following ileocolonic resection for Crohn's disease—considerations in pre-operative and peri-operative optimisation techniques to improve outcome. *J Crohns Colitis*. 2015;9:483–491.
135. Zuo L, Li Y, Wang H, et al. A practical predictive index for intra-abdominal septic complications after primary anastomosis for Crohn's disease: change in C-reactive protein level before surgery. *Dis Colon Rectum*. 2015;58:775–781.
136. Li Y, Stocchi L, Cherla D, et al. Association of preoperative narcotic use with postoperative complications and prolonged length of hospital stay in patients with Crohn disease. *JAMA Surg*. 2016;151:726–734.
137. Germain A, Guéant RM, Chamaillard M, et al. NOD2 gene variant is a risk factor for postoperative complications in patients with Crohn's disease: a genetic association study. *Surgery*. 2016;160:74–80.
138. Yamamoto T, Spinelli A, Suzuki Y, et al. Risk factors for complications after ileocolonic resection for Crohn's disease with a major focus on the impact of pre-operative immunosuppressive and biologic therapy: a retrospective international multicentre study. *United Eur Gastroenterol J*. 2016;4:784–793.
139. Sahami S, Bartels SA, D'Hoore A, et al. A multicentre evaluation of risk factors for anastomotic leakage after restorative proctocolectomy with ileal pouch-anal anastomosis for inflammatory bowel disease. *J Crohns Colitis*. 2016;10:773–778.
140. Guo K, Ren J, Li G, et al. Risk factors of surgical site infections in patients with Crohn's disease complicated with gastrointestinal fistula. *Int J Colorectal Dis*. 2017;32:635–643.
141. Diederer K, Sahami SS, Tabbers MM, et al. Outcome after restorative proctocolectomy and ileal pouch-anal anastomosis in children and adults. *Br J Surg*. 2017;104:1640–1647.
142. Heimann TM, Swaminathan S, Greenstein AJ, et al. Incidence and factors correlating with incisional hernia following open bowel resection in patients with inflammatory bowel disease: a review of 1000 patients. *Ann Surg*. 2018;267:532–536.
143. Huang W, Tang Y, Nong L, et al. Risk factors for postoperative intra-abdominal septic complications after surgery in Crohn's disease: a meta-analysis of observational studies. *J Crohns Colitis*. 2015;9:293–301.
144. Beddy D, Dozois EJ, Pemberton JH. Perioperative complications in inflammatory bowel disease. *Inflamm Bowel Dis*. 2011;17:1610–1619.
145. Luong TV, Grandt SD, Negoï I, Palubinskas S, El-Hussuna A. Preoperative factors associated with prolonged postoperative in-hospital length of stay in patients with Crohn's disease undergoing intestinal resection or strictureplasty. *Int J Colorectal Dis*. 2019;34:1925–1931. doi:10.1007/s00384-019-03418-8