

Living with ulcerative colitis study (LUCY) in England: A retrospective study evaluating healthcare resource utilisation and direct healthcare costs of postoperative care in ulcerative colitis

Item Type	Journal article
Authors	Brookes, Matthew;Waller, J;Cappelleri, JC;Modesto, I;Dibonaventura, MD;Bohm, N;Mokgokong, R;Massey, O;Wood, R;Bargo, D
Citation	Brookes, M.J., Waller, J., Cappelleri, J.C., et al. (2020) Living with Ulcerative Colitis Study (LUCY) in England: a retrospective study evaluating healthcare resource utilisation and direct healthcare costs of postoperative care in ulcerative colitis. <i>BMJ Open Gastroenterology</i> , 7:e000456. doi:10.1136/ bmjgast-2020-000456
DOI	10.1136/bmjgast-2020-000456
Publisher	BMJ
Journal	BMJ Open Gastroenterology
Rights	Licence for published version: Creative Commons Attribution-NonCommercial 4.0 International
Download date	2026-06-18 14:54:51
License	http://creativecommons.org/licenses/by-nc/4.0/
License	http://creativecommons.org/licenses/by-nc/4.0/
Link to Item	http://hdl.handle.net/2436/624722

Living with Ulcerative Colitis Study (LUCY) in England: a retrospective study evaluating healthcare resource utilisation and direct healthcare costs of postoperative care in ulcerative colitis

Matthew J Brookes ,^{1,2} John Waller,³ Joseph C Cappelleri,⁴ Irene Modesto,⁵ Marco D DiBonaventura,⁶ Natalie Bohm,⁷ Ruth Mokgokong,⁷ Olivia Massey,³ Robert Wood,³ Danielle Bargo⁶

To cite: Brookes MJ, Waller J, Cappelleri JC, *et al.* Living with Ulcerative Colitis Study (LUCY) in England: a retrospective study evaluating healthcare resource utilisation and direct healthcare costs of postoperative care in ulcerative colitis. *BMJ Open Gastro* 2020;**7**:e000456. doi:10.1136/bmjgast-2020-000456

► Additional material is published online only. To view please visit the journal online (<http://dx.doi.org/10.1136/bmjgast-2020-000456>).

Received 29 May 2020
Revised 15 July 2020
Accepted 17 July 2020



© Author(s) (or their employer(s)) 2020. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Dr Marco D DiBonaventura;
Marco.DiBonaventura@pfizer.com

ABSTRACT

Objective Ulcerative colitis (UC) is a lifelong, relapsing-remitting disease. Patients non-responsive to pharmacological treatment may require a colectomy. We estimated pre-colectomy and post-colectomy healthcare resource utilisation (HCRU) and costs in England.

Design/Method A retrospective, longitudinal cohort study indexing adult patients with UC undergoing colectomy (2009–2015), using linked Clinical Practice Research Datalink/Hospital Episode Statistics data, was conducted. HCRU, healthcare costs and pharmacological treatments were evaluated during 12 months prior to and including colectomy (baseline) and 24 months post-colectomy (follow-up; F-U), comparing baseline/F-U, emergency/elective colectomy and subtotal/full colectomy using descriptive statistics and paired/unpaired tests.

Results 249 patients from 26 165 identified were analysed including 145 (58%) elective and 184 (74%) full colectomies. Number/cost of general practitioner consultations increased post-colectomy ($p < 0.001$), and then decreased at 13–24 months ($p < 0.05$). From baseline to F-U, the number of outpatient visits, number/cost of hospitalisations and total direct healthcare costs decreased (all $p < 0.01$). Postoperative HCRU was similar between elective and emergency colectomies, except for the costs of colectomy-related hospitalisations and medication, which were lower in the elective group ($p < 0.05$). Postoperative costs were higher for subtotal versus full colectomies ($p < 0.001$). At 1–12 month F-U, 30%, 19% and 5% of patients received aminosaliclates, steroids and immunosuppressants, respectively.

Conclusion HCRU/costs increased for primary care in the first year post-colectomy but decreased for secondary care, and varied according to the colectomy type. Ongoing and potentially unnecessary pharmacological therapy was seen in up to 30% of patients. These findings can inform patients and decision-makers of potential benefits and burdens of colectomy in UC.

INTRODUCTION

Ulcerative colitis (UC) is a relapsing-remitting inflammatory bowel disease.^{1–3} Patients usually present with bloody diarrhoea, abdominal pain,

Key messages

What is already known about this subject?

- Despite numerous pharmacological therapy options for the management of ulcerative colitis (UC), some patients fail to respond to these and require colectomy.
- Although active UC can impact health-related quality of life and healthcare resource utilisation (HCRU), post-colectomy patients should not be considered free from these burdens.
- Approximately one-third of patients having UC-related surgery experience some form of postoperative complications.
- Studies report high costs associated with UC-related surgery; however, there is a paucity of published data on HCRU in the postoperative setting in the UK.

What are the new findings?

- Our study showed changes in HCRU and costs during perioperative and postoperative periods in patients with UC undergoing colectomy in the UK.
- We found that after surgery, HCRU changed, with increasing usage of primary care in the first 12 months after surgery.
- We also demonstrated that predominant HCRU is similar in patients undergoing emergency versus elective colectomies; however, HCRU was significantly higher in subtotal colectomy patients for emergency versus elective procedures.

urgency and tenesmus.^{1–4} Globally, UC prevalence ranges from 2.4 to 505 cases/100 000 people, with an estimated annual incidence of 10/100 000 people, and a prevalence of approximately 240/100 000 (around 146 000 cases) in the UK.^{5,6}

Various pharmacological UC treatments exist, including aminosaliclates (5-ASAs), corticosteroids, immunosuppressants, biologics



Key messages

How might it impact on clinical practice in the foreseeable future?

- ▶ Our findings will enable better preoperative planning and counselling for patients due to have a colectomy.
- ▶ Engaging with the primary care sector and/or physicians may reduce the potentially unnecessary pharmacological therapies during the postoperative period, with some patients still receiving therapies at 24 months post-colectomy.
- ▶ Our study may enable the development of a postoperative toolkit resource for patients and their healthcare professionals, to outline a defined postoperative management plan and provide patients with information on what to expect.
- ▶ Quantifying the relationship between colectomy and costs will also inform future health economic studies.

and tofacitinib. European guidelines recommend sustained steroid-free remission as the treatment goal.^{7–9} Despite these options, some patients with UC require a colectomy.¹⁰

The highest colectomy rates are seen during the first few years after diagnosis.¹¹ Patients usually undergo colectomy due to acute severe UC (10%–60% of patients).^{12–14} A retrospective analysis reported that 56% of patients with UC with a primary non-response to infliximab underwent colectomy.¹⁵ International studies reported colectomies in 7%–10% of patients with UC, with decreasing colectomy rates over recent years.^{16–18}

Active UC is associated with reduced health-related quality of life (HRQoL) and productivity,^{19–24} frequent visits to accident and emergency departments (A&E), hospital stays and healthcare costs,^{24–26} with greater costs with increasing frequency of relapse.²⁷ Although colectomy may be life-saving, it was recommended not to be considered a cure for UC.²⁸ Patients undergoing colectomy can still experience poor HRQoL,^{29–31} anxiety and depression, problems with body image and sexual function, and decreased productivity.³² Approximately one-third of patients having UC-related surgery experience some form of postoperative complications³³; and these can result in substantial humanistic and economic burden.³⁴

Although studies report high costs associated with UC-related surgery, there are limited data on healthcare resource utilisation (HCRU) in postoperative settings.^{35 36} Surgery in UC may be an emergency or elective procedure³⁷; costs associated with emergency versus elective procedures were reported to be higher.³⁸

We aimed to estimate preoperative and postoperative HCRU and direct healthcare costs among patients with UC undergoing a colectomy, and to examine preoperative and postoperative patterns of concomitant medication use, postoperative complications and the association of colectomy type with HCRU, direct healthcare costs and postoperative medication use.

MATERIALS AND METHODS

Study design

This study used linked Clinical Practice Research Datalink (CPRD) GOLD and Hospital Episode Statistics (HES) data (this study is based in part on data from the Clinical Practice Research Datalink obtained under licence from the UK Medicines and Healthcare products Regulatory Agency. The data are provided by patients and collected by the NHS as part of their care and support. The interpretation and conclusions contained in this study are those of the author/s alone; copyright 2017, reused with the permission of the Health and Social Care Information Centre. All rights reserved; the OPCS Classification of Interventions and Procedures, codes, terms and text is Crown copyright (2016) published by the Health and Social Care Information Centre, also known as NHS Digital, and licensed under the Open Government Licence available at <http://www.nationalarchives.gov.uk/doc/open-government-licence/version/3/>). CPRD captures anonymised data related to all primary care patient interactions at participating general practitioner (GP) practices in the UK.³⁹ HES captures details on all secondary care patient interactions at NHS hospitals in England, with limited data on treatments prescribed; only high-cost drugs are observed and specific treatments can rarely be identified.⁴⁰

A retrospective, longitudinal cohort study was conducted (online supplementary figure 1). The study cohort comprised of adult patients (aged 18 or over) undergoing a colectomy between 1 January 2009 and 31 December 2015 (indexing period), with a prior UC diagnosis, and continuously registered with the GP practice for the 12-month period prior to (ie, baseline) and 24-month period following (ie, follow-up; F-U) colectomy. The first/earliest colectomy observed determined the index date. Patients with a colectomy recorded in the 12 months prior to the index event were excluded, as were patients with a diagnosis of Crohn's disease and/or cancer (due to overlap in high-cost drug coding for UC and cancer treatments) recorded within the study period (1 January 2008–31 December 2017). Code lists for study cohort identification are detailed in online supplementary table S1.

Patients undergoing elective and emergency colectomies (defined according to type of hospital admission), and subtotal and full colectomies, were considered. As the latter could not be ascertained from the procedural coding system used, the index colectomy was classified as subtotal if an additional colectomy-related procedure was recorded during F-U.

Outcomes

Outcomes measured included baseline demographics and clinical characteristics, HCRU/costs and postoperative complications.

Direct healthcare costs were calculated at baseline and F-U by imputing unit costs to a range of healthcare resources. Costs of prescriptions (primary care only) for

5-ASAs, steroids and immunosuppressants were calculated using net ingredient costs from Prescription Cost Analysis 2017 tables.⁴¹ GP consultation costs were calculated by applying unit costs from the Personal Social Services Research Unit.⁴² Costs of hospital interactions were calculated by deriving Healthcare Resource Groups (HRGs) for each visit/stay and subsequently applying national tariffs (2018/2019) for each HRG.⁴³ Costs pertaining to A&E visits resulting in hospitalisation and additional procedures/surgeries are incorporated into HRGs derived for each visit/stay; therefore, specific costs relating to these resources cannot be identified. Costs of stoma care were not included due to data limitations.

Costs associated with the index colectomy (and the inpatient spell during which the index colectomy was performed) were attributed to the baseline period as the hospital admission date will have occurred prior to the index date, that is, date of colectomy, by definition.

Postoperative complications were assessed (online supplementary table S2) via the presence of diagnostic codes recorded following index colectomy, with gastrointestinal (GI) complications assessed throughout F-U and all other complications in the first 30 days of F-U only (widened to 60 days in a sensitivity analysis).

Due to coding limitations for high-cost drugs in HES data, specific biologic therapies prescribed/administered could not be identified; therefore, use of biologic therapies could not be assessed in this study.

Analysis

All outcomes were analysed descriptively and reported using frequencies and percentages for categorical, and counts, means, medians and SD for continuous variables.

Significance testing between baseline and F-U was conducted using paired t-tests for numeric and McNemar's test for dichotomous outcomes. Significance testing between patient subgroups was conducted using t-tests for numeric and χ^2 or Fisher's exact tests (if any expected cell count was ≤ 5) for dichotomous outcomes.

All statistical analyses were performed using Stata V.15.1 or later.⁴⁴

Ethics

The study was approved by an Independent Scientific Advisory Committee (Protocol No.: 18_263).

RESULTS

Eligible patients

In total, 26 165 patients were indexed; 249 patients had a prior UC diagnosis and satisfied selection criteria. Numbers of patients indexed decreased from 48 (2009) to 10 (2015). Reasons for patient exclusion are in online supplementary figure 2.

Demographics and disease characteristics

Demographic and clinical characteristics are shown in table 1.

Table 1 Baseline demographics and disease characteristics

	Total
	N=249
Age (years)	
Mean (SD)	50.5 (16.6)
Min to max	19 to 86
Median (IQR)	51 (36–64)
Sex, n (%)	
Female	105 (42%)
Male	144 (58%)
BMI (kg/m ²)	
N	233
Mean (SD)	26.6 (5.7)
Min to max	15.7 to 55.7
Median (IQR)	25.7 (22.5–29.7)
Age at diagnosis (years)	
Mean (SD)	46.6 (16.6)
Min to max	10 to 86
Median (IQR)	47 (33–60)
Time since diagnosis (years)	
Mean (SD)	3.8 (4.0)
Min to max	0 to 17.3
Median (IQR)	2.03 (0.6–6.6)
Categorised, n (%)	
≤ 1 year	82 (33%)
> 1 year	167 (67%)
CCI	
Mean (SD)	0.6 (1.3)
Min to max	0 to 8
Median (IQR)	0 (0–1)
Categorised, n (%)	
0	181 (73%)
1–2	51 (21%)
3–4	17 (7%)
5+	0 (0%)
Disease extent, n (%)	
Pancolitis	38 (15%)
Left-sided	11 (4%)
Proctosigmoiditis	56 (23%)
Proctitis	89 (36%)
Other/Unknown	55 (22%)
Index colectomy (top five procedures only)*, n (%)	
Other specified subtotal excision of colon (H29.8)	37 (15%)
Colectomy and ileostomy NEC (H11.4)	35 (14%)

Continued

Table 1 Continued

	Total
Total colectomy and ileostomy NEC (H05.3)	32 (13%)
Panproctocolectomy and ileostomy (H04.1)	29 (12%)
Unspecified subtotal excision of colon (H29.9)	20 (8%)
Emergency/elective colectomy, n (%)	
Emergency	104 (42%)
Elective	145 (58%)
Subtotal/full colectomy†, n (%)	
Subtotal	65 (26%)
Full	184 (74%)

*All other colectomy-related procedures (n=28 in total) at index were performed in <6% of patients.

†Determined based on the number of colectomy-related procedure codes observed within the study period (1=full, ie, index only; >1=subtotal, ie, index and subsequent procedure). BMI, body mass index; CCI, Charlson Comorbidity Index.

HCRU and direct healthcare costs

HCRU and costs at baseline and F-U

GP consultations increased post-colectomy from baseline to 1–12 month F-U ($p<0.001$; [figure 1](#)). Hospital stays, outpatient visits and hospital admissions via A&E decreased post-colectomy from baseline to 1–12 month F-U ($p<0.01$; [figure 1](#)). GP consultations, outpatient visits and hospital admissions via A&E post-colectomy decreased at 13–24 month versus 1–12 month F-U ($p<0.001$; [figure 1](#)). Each HCRU element was lower at 13–24 month F-U versus baseline ($p<0.05$; [figure 1](#)).

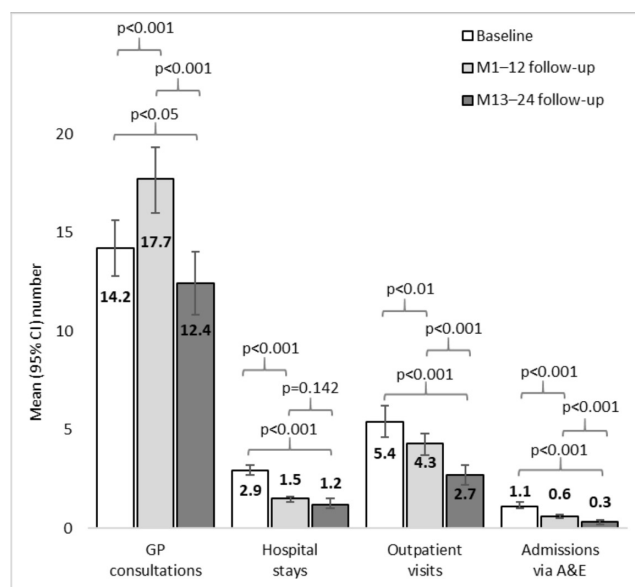


Figure 1 Healthcare resource utilisation at baseline and follow-up. Error bars represent 95% CIs. A&E, accident and emergency; GP, general practitioner; M, month.

Although outpatient visits decreased following surgery, mean number of visits to the colorectal surgery department increased from 0.9 at baseline to 2.1 at 1–12 month F-U ($p<0.001$), and then decreased to 1.2 at 13–24 month F-U ($p<0.001$). The mean length of individual hospital stay decreased from 12.6 days at baseline to 3.4 days at 1–12 month F-U ($p<0.001$) and 2.4 days at 13–24 month F-U ($p<0.05$ vs 1–12 month F-U; $p<0.001$ vs baseline).

Mean total direct healthcare costs at baseline, 1–12 month and 13–24 month F-U were £10366, £4433 and £3333, respectively, with higher costs at baseline driven by costs of the inpatient spell needed for index colectomy. Costs were lower at 1–12 month F-U versus baseline ($p<0.001$; [figure 2A](#)).

GP consultation costs increased post-colectomy from baseline to 1–12 month F-U ($p<0.001$). Hospital stay costs related to colectomy and medication decreased from baseline to 1–12 month F-U ($p<0.001$), while costs of outpatient visits and hospital stays unrelated to colectomy were similar at baseline and 1–12 month F-U ([figure 2A](#)). Costs of GP consultations, outpatient visits, hospital stays unrelated to colectomy and medication (5-ASA, steroid and immunosuppressant therapies only) decreased at 13–24 month vs 1–12 month F-U, with changes statistically significant except for costs of medication ($p<0.01$; [figure 2A](#)). Costs of hospital stays related to colectomy remained similar at 13–24 month and 1–12 month F-U ([figure 2A](#)). Costs of GP consultations were comparable between baseline and 13–24 month F-U. All other costs were lower at 13–24 month F-U versus baseline (all $p<0.01$).

HCRU and costs by emergency/elective colectomy

At baseline, numbers of hospitalisations and outpatient visits were higher and admissions via A&E were lower in the elective versus emergency cohort ($p<0.05$). The length of hospital stay from admission to colectomy and from colectomy to discharge was lower in elective versus emergency colectomy patients ($p<0.01$). Otherwise, there were limited differences in HCRU during F-U.

Mean total direct healthcare costs in emergency and elective colectomy patients were comparable at baseline (£10133 and £10533, respectively) and numerically higher for emergency colectomy patients at 24 month F-U (£8595 and £7171, respectively). At baseline, costs of outpatient visits and medication were higher in elective versus emergency colectomy patients ($p<0.001$). F-U costs of hospital stays related to colectomy and medication were lower in elective versus emergency colectomy patients ($p<0.05$). All other costs were comparable at baseline and F-U between elective versus emergency colectomy patients.

HCRU and costs by subtotal/full colectomy

At baseline, number of outpatient visits was higher in subtotal versus full colectomy patients ($p<0.05$), but otherwise there were limited differences in HCRU. During F-U,

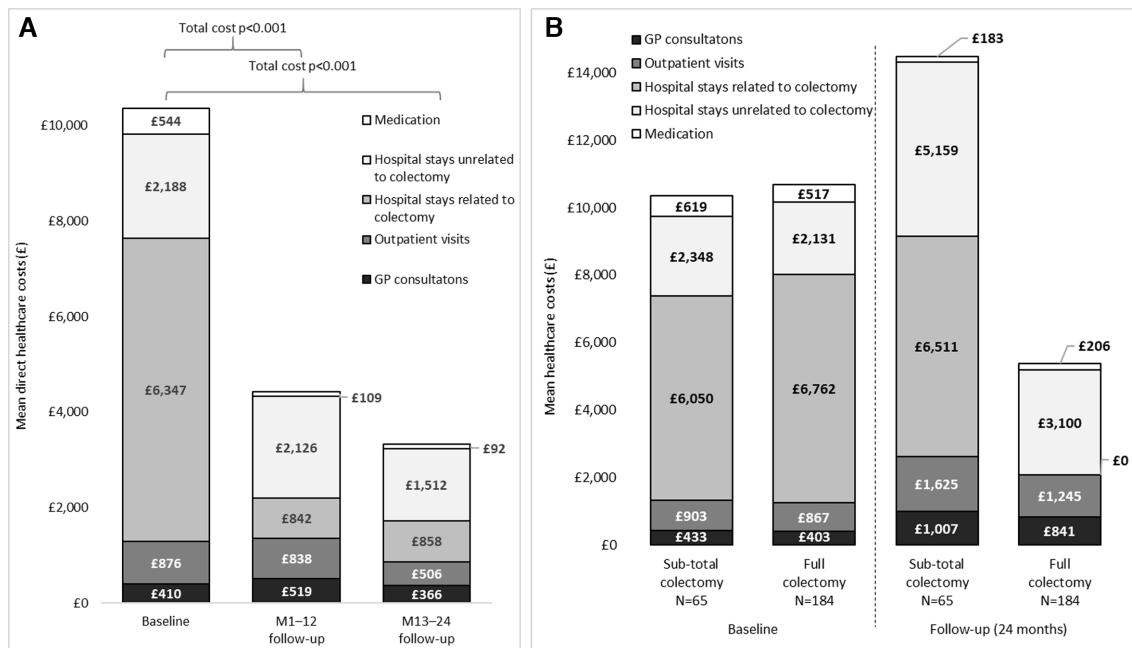


Figure 2 Direct healthcare costs (A) during baseline and follow-up; and (B) by subtotal/full colectomy. (A) The cost of the index colectomy is included in baseline costs. Medication costs include 5-ASAs, steroids and immunosuppressants. (B) Medication costs include 5-ASAs, steroids and immunosuppressants. 5-ASAs, aminosaliculates; GP, general practitioner; M, month.

GP consultations were comparable between subtotal and full colectomy patients; however, the number of outpatient visits, and number and length of hospital stays, were greater in subtotal versus full colectomy patients ($p < 0.0001$), as was the proportion of patients prescribed 5-ASAs ($p < 0.05$).

There were no differences in direct costs in full versus subtotal colectomy patients at baseline (figure 2B). During F-U, total direct costs were higher in subtotal versus full colectomy patients ($p < 0.0001$; figure 2B). Direct costs of GP consultations, hospital stays (both colectomy-related and non-colectomy-related) and outpatient visits were higher in subtotal versus full colectomy patients ($p < 0.05$; figure 2B).

Pharmacological treatment

Although proportions of patients receiving 5-ASAs, steroids and immunosuppressants were lower at 1–12 month F-U versus baseline ($p < 0.001$), a substantial proportion of patients received treatment with 5-ASAs

(30%), steroids (19%) and/or immunosuppressants (5%) in 12 months following index colectomy. Proportions of patients receiving 5-ASAs and steroids decreased from 1 to 12 month to 13–24 month F-U ($p < 0.01$), while immunosuppressant use remained stable across two F-U periods (figure 3A). The most commonly prescribed medications in each class during both baseline and F-U were mesalazine (5-ASA), prednisolone and azathioprine.

The proportion of patients receiving 5-ASAs, steroids and immunosuppressants at baseline was numerically higher in elective versus emergency colectomy patients ($p < 0.001$ for immunosuppressants). Conversely, the proportion of patients receiving 5-ASAs and steroids post-colectomy was numerically lower in elective versus emergency colectomy patients ($p < 0.05$ for 5-ASAs).

A higher proportion of subtotal versus full colectomy patients received 5-ASA (77% vs 61%, respectively; $p < 0.05$) and steroid treatment (69% vs 53%, respectively; $p < 0.05$) at baseline, while immunosuppressant use

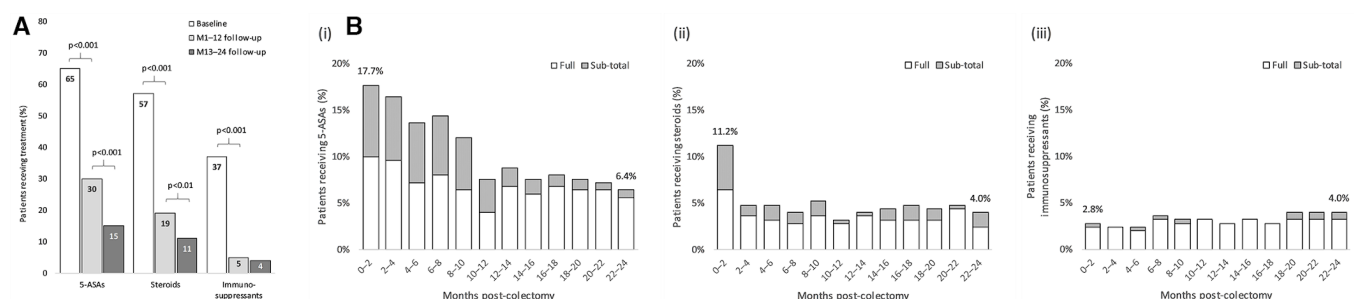


Figure 3 Pharmacological treatment at (A) baseline and follow-up; and (B) follow-up (i) 5-ASAs, (ii) steroids and (iii) immunosuppressants. 5-ASAs, aminosaliculates; M, month.



was numerically higher in subtotal versus full colectomy patients (45% vs 34%, respectively). At 1–12 month F-U after the initial colectomy, a higher proportion of subtotal versus full colectomy patients received 5-ASA treatment (59% vs 20%, respectively; $p < 0.001$), while a numerically higher proportion of subtotal versus full colectomy patients received steroid treatment (26% vs 16%, respectively), and administration of immunosuppressants was comparable in subtotal and full colectomy patients.

Use of 5-ASAs decreased in the first 10 months following colectomy and then stabilised, while steroid use decreased after the first 2 months of F-U to a constant level (figure 3B). The proportion of patients receiving immunosuppressants remained stable from immediately after the index date until end of F-U at 24 months (figure 3B).

Postoperative complications

During F-U, 39% of patients experienced postoperative GI complications, with 21% of these suspected of experiencing chitinous (ICD-10: K91.8—Other postprocedural disorders of digestive system, not elsewhere classified). Steroid use at baseline was not associated with postoperative GI complications, which were experienced by 38% of patients with no steroid use and 40% of patients with steroids ($p = 0.795$).

Fewer than 10% of patients reported any other type of complication during the 30-day postoperative period; widening the observation period to 60 days in a sensitivity analysis did not impact findings.

The proportion of patients experiencing postoperative complications was similar in patients having emergency and elective surgery.

DISCUSSION

This analysis of primary and secondary care data from the UK explored HCRU, healthcare costs and medication use in patients with UC prior to and following colectomy, and the potential influence of emergency versus elective and subtotal versus full colectomy on postoperative HCRU, costs and medication. We found a lower annual colectomy rate between January 2009 and December 2015 than that reported in another HES data study between April 1997 and March 2012.⁴⁵ However, this could be explained by the decrease in colectomy rates over time.^{16 46 47}

Visits to the colorectal surgery department and GP consultations increased during F-U, probably reflecting increased patient monitoring after surgery; however, use of most secondary care resources decreased following colectomy. Although direct healthcare costs decreased from baseline (mean cost/patient $>£10\,000$), they remained substantial 12 months later (mean cost/patient $>£4000$). Mean total costs associated with UC-related surgery over 6 months in a UK study were reported to be approximately $£15\,000$,³⁵ while in the US, colectomy and 6 months follow-up costs were $\$90\,445$.³⁶

Despite the total direct healthcare costs at baseline being similar, regardless of whether colectomy was elective or emergency, the mean total direct healthcare costs during F-U were 17% lower in patients undergoing elective colectomy, yet statistical significance was not observed—likely a result of the limited sample size studied. Previous studies have shown higher cost differences but in different healthcare systems. A Canadian study reported a two-thirds higher cost in emergency versus elective colectomies.³⁸ A US National Inpatient Sample also suggested that costs and outcomes of emergency colectomies depended on whether the surgery occurred within the first 24 hours, with greater complications and costs if colectomies were delayed until >24 hours after admission.⁴⁸ In our analysis, the mean time from admission to colectomy was 1.3 days in elective vs 10.5 days in emergency colectomies.

We showed that although use of 5-ASAs, steroids and immunosuppressants decreased following surgery, around one-third and one-fifth of patients, respectively, received 5-ASAs and steroids even after surgery, with these increasing to three-fifths and one-third, respectively, when subtotal colectomy patients were considered. This supports the view that colectomy is not a cure for UC and even patients undergoing full colectomy can require further medical or surgical care.^{28 33} However, it should be noted that the indication of prescribed medications is not captured in these data, but despite this, a substantial proportion of patients still appear to receive 5-ASA, immunosuppressant and steroid treatment following full colectomy. Further research is required to fully understand this.

Almost 40% of patients experienced GI complications at 24 month F-U, but the incidence of other postoperative complications was very low. No differences were seen in our analysis between elective and emergency surgery in rates of any postoperative complications, which is similar to a US tertiary care study which reported complications in 47% of patients undergoing colectomy.⁴⁹ By contrast, a Canadian study reported postoperative complications in only 27% of patients, with a significantly higher risk of post-colectomy complications in emergency surgery patients.⁵⁰ A systematic review identified a high level of variability in rates of postoperative complications; those up to 30 days postoperatively occurring in 9%–65% of patients and those after 30 days postoperatively occurring in 17%–55% of patients.³³ However, it should be noted that due to coding limitations within HES, it was not possible to explicitly identify complications due to colectomy. Instead, the presence of diagnostic codes indicative of complications typically related to colectomy were used.

This study had limitations. Linkage of the CPRD to HES reduced the sample from the UK to English patients, and patients receiving private medical care, in prisons, some residential homes or homeless were not represented. Identifying patients with UC via ICD-10 and Read diagnosis codes might have led to inappropriate patients' inclusion/exclusion. Overestimation of HCRU/

costs might have occurred by including resource use for non-UC-related conditions. Underestimation of HCRU/costs might have resulted from excluding advanced therapy and stoma care costs; lack of national tariff for some HRGs (some prices negotiated locally); and HRGs derived using the most recent Local Payment Group, possibly underestimating costs of earlier admissions (costs of technology/medications decreased over time).

In conclusion, although HCRU and healthcare costs in patients with UC decreased post-colectomy, there was still a substantial HCRU and economic burden postoperatively. Patients undergoing colectomy still require medical attention and use healthcare resources, which should be considered in their follow-up.

Author affiliations

¹Gastroenterology Department, Royal Wolverhampton NHS Trust, Wolverhampton, UK

²Research Institute in Healthcare Science, Faculty of Science and Engineering, University of Wolverhampton, Wolverhampton, UK

³Adelphi Real World, Bollington, UK

⁴Biostatistics, Pfizer Inc, Groton, Connecticut, USA

⁵Inflammation & Immunology, Pfizer Inc, New York, New York, USA

⁶Patient Health Impact, Pfizer Inc, New York, New York, USA

⁷Pfizer Inc, Tadworth, UK

Acknowledgements Medical writing support, under the guidance of the authors, was provided by Carole Evans, PhD, on behalf of Adelphi Real World, and was funded by Pfizer Inc in accordance with Good Publication Practice (GPP3) guidelines (Ann Intern Med 2015;163:461–464).

Contributors JCC, MDD, NB, RM and DB were involved in the conception and design of the study, analysis and interpretation of data, revising the article critically for important intellectual content, and final approval of the version to be submitted. MJB and IM were involved in the analysis and interpretation of data, revising the article critically for important intellectual content, and final approval of the version to be submitted. JW, OM and RW were involved in the conception and design of the study, acquisition of data, analysis and interpretation of data, drafting the article and revising it critically for important intellectual content, and final approval of the version to be submitted.

Funding The study was funded by Pfizer Inc.

Disclaimer The research department of MJB has received grant support from Tillotts Pharma and Vifor Pharma (Switzerland). MJB has received honoraria and travel support for consulting or lecturing from Vifor Pharma, Tillotts Pharma and AbbVie. DB, JCC, IM, MDD and RM are employees and stockholders of Pfizer Inc (which funded this study). NB is an employee of Pfizer Inc and stockholder of Pfizer Inc (which funded this study). JW, OM and RW are employed by Adelphi Real World, which received funding from Pfizer Inc in connection with the development of this manuscript.

Competing interests The research department of MJB has received grant support from Tillotts Pharma and Vifor Pharma (Switzerland). MJB has received honoraria and travel support for consulting or lecturing from Vifor Pharma, Tillotts Pharma and AbbVie. DB, JCC, IM, MDD and RM are employees and stockholders of Pfizer Inc (which funded this study). NB is an employee of Pfizer Inc and stockholder of Pfizer Inc (which funded this study). JW, OM and RW are employed by Adelphi Real World, which received funding from Pfizer Inc in connection with the development of this manuscript.

Patient consent for publication Not required.

Ethics approval This was a retrospective analysis of data, involving no decisions regarding patient interventions or the omission of interventions, and all patient-level data in the data sources were anonymised. Hence, institutional review board/ethics approval and patient informed consent were not needed. Use of linked CPRD-HES data required Independent Scientific Advisory Committee approval; an abridged version of the protocol was supplied to the committee and approval was granted (Protocol No.: 18_263).

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data may be obtained from a third party and are not publicly available. The deidentified patient data accessed and analysed for the purposes of this study are available from the Clinical Practice Research Datalink (email: enquiries@cprd.com) and access to these data are permissible on approval of a written study protocol by the Independent Scientific Advisory Committee.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iD

Matthew J Brookes <http://orcid.org/0000-0002-8782-0292>

REFERENCES

- 1 Ungaro R, Mehandru S, Allen PB, *et al*. Ulcerative colitis. *Lancet* 2017;389:1756–70.
- 2 Antonelli E, Villanacci V, Bassotti G. Novel oral-targeted therapies for mucosal healing in ulcerative colitis. *World J Gastroenterol* 2018;24:5322–30.
- 3 Satsangi J, Silverberg MS, Vermeire S, *et al*. The Montreal classification of inflammatory bowel disease: controversies, consensus, and implications. *Gut* 2006;55:749–53.
- 4 Feuerstein JD, Cheifetz AS. Ulcerative colitis: epidemiology, diagnosis, and management. *Mayo Clin Proc* 2014;89:1553–63.
- 5 Ng SC, Shi HY, Hamidi N, *et al*. Worldwide incidence and prevalence of inflammatory bowel disease in the 21st century: a systematic review of population-based studies. *Lancet* 2018;390:2769–78.
- 6 Ulcerative colitis: management. National Institute for Health and Care Excellence, 2013. Available: <https://www.nice.org.uk/guidance/cg166> [Accessed 17 Apr 2019].
- 7 Danese S, Allez M, van Bodegraven AA, *et al*. Unmet medical needs in ulcerative colitis: an expert group consensus. *Dig Dis* 2019;37:266–83.
- 8 Wehkamp J, Stange EF. Recent advances and emerging therapies in the non-surgical management of ulcerative colitis [version 1; peer review: 3 approved]. *F1000Res* 2018;7:1207.
- 9 Harbord M, Eliakim R, *et al*. Third European evidence-based consensus on diagnosis and management of ulcerative colitis. Part 2: current management. *J Crohns Colitis* 2017;11:769–84.
- 10 Wong DJ, Roth EM, Feuerstein JD, *et al*. Surgery in the age of biologics. *Gastroenterol Rep* 2019;7:77–90.
- 11 Solberg IC, Lygren I, *et al*, IBSEN Study Group. Clinical course during the first 10 years of ulcerative colitis: results from a population-based inception cohort (IBSEN study). *Scand J Gastroenterol* 2009;44:431–40.
- 12 Lynch RW, Churchhouse AMD, *et al*, UK IBD Audit Steering Group. Predicting outcome in acute severe ulcerative colitis: comparison of the Travis and HO scores using UK IBD audit data. *Aliment Pharmacol Ther* 2016;43:1132–41.
- 13 NICE clinical guideline and quality standard: ulcerative colitis scope. National Institute for Health and Care Excellence, 2019. Available: <https://www.nice.org.uk/guidance/cg166/documents/ulcerative-colitis-final-scope2> [Accessed 17 Apr 2019].
- 14 Turner D, Walsh CM, Steinhart AH, *et al*. Response to corticosteroids in severe ulcerative colitis: a systematic review of the literature and a meta-regression. *Clin Gastroenterol Hepatol* 2007;5:103–10.
- 15 Papamichael K, Rivals-Lerebours O, Billiet T, *et al*. Long-term outcome of patients with ulcerative colitis and primary non-response to infliximab. *J Crohns Colitis* 2016;10:1015–23.
- 16 Parragi L, Fournier N, *et al*, Swiss IBD Cohort Study Group. Colectomy rates in ulcerative colitis are low and decreasing: 10-year follow-up data from the Swiss IBD Cohort Study. *J Crohns Colitis* 2018;12:811–8.
- 17 Manetti N, Bagnoli S, Rogai F, *et al*. Disease course and colectomy rate of ulcerative colitis: a follow-up cohort study of a referral center in Tuscany. *Inflamm Bowel Dis* 2016;22:1945–53.
- 18 Targownik LE, Singh H, Nugent Z, *et al*. The epidemiology of colectomy in ulcerative colitis: results from a population-based cohort. *Am J Gastroenterol* 2012;107:1228–35.
- 19 Yaras A, Rubin DT, Panés J, *et al*. Burden of ulcerative colitis on functioning and well-being: a systematic literature review of the SF-36® health survey. *J Crohns Colitis* 2018;12:600–9.
- 20 Yaras A, Maher SM, Bayliss MS, *et al*. Psychometric validation of the work productivity and activity impairment questionnaire in



- ulcerative colitis: results from a systematic literature review. *J Patient Rep Outcomes* 2018;2:62.
- 21 Van Assche G, Peyrin-Biroulet L, Sturm A, *et al.* Burden of disease and patient-reported outcomes in patients with moderate to severe ulcerative colitis in the last 12 months - Multicenter European cohort study. *Dig Liver Dis* 2016;48:592–600.
 - 22 Vaizey CJ, Gibson PR, Black CM, *et al.* Disease status, patient quality of life and healthcare resource use for ulcerative colitis in the UK: an observational study. *Frontline Gastroenterol* 2014;5:183–9.
 - 23 Parra RS, Chebli JMF, Amarante HMBS, *et al.* Quality of life, work productivity impairment and healthcare resources in inflammatory bowel diseases in Brazil. *World J Gastroenterol* 2019;25:5862–82.
 - 24 Gibson PR, Vaizey C, Black CM, *et al.* Relationship between disease severity and quality of life and assessment of health care utilization and cost for ulcerative colitis in Australia: a cross-sectional, observational study. *J Crohns Colitis* 2014;8:598–606.
 - 25 Cohen R, Skup M, Ozbay AB, *et al.* Direct and indirect healthcare resource utilization and costs associated with ulcerative colitis in a privately-insured employed population in the US. *J Med Econ* 2015;18:447–56.
 - 26 Hudesman DP, Chakravarty SD, Emond B, *et al.* Healthcare resource utilization and costs associated with inflammatory bowel disease among patients with chronic inflammatory diseases: a retrospective cohort study. *BMC Rheumatol* 2020;4:16.
 - 27 Bodger K, Yen L, Szende A, *et al.* Medical resource utilization and associated costs in patients with ulcerative colitis in the UK: a chart review analysis. *Eur J Gastroenterol Hepatol* 2014;26:213–21.
 - 28 Dayan B, Turner D. Role of surgery in severe ulcerative colitis in the era of medical rescue therapy. *World J Gastroenterol* 2012;18:3833–8.
 - 29 Abolfotouh S, Rautio T, Klintrup K, *et al.* Predictors of quality-of-life after ileal pouch-anal anastomosis in patients with ulcerative colitis. *Scand J Gastroenterol* 2017;52:1078–85.
 - 30 Koerdt S, Jehle EC, Kreis ME, *et al.* Quality of life after proctocolectomy and ileal pouch-anal anastomosis in patients with ulcerative colitis. *Int J Colorectal Dis* 2014;29:545–54.
 - 31 Raviram S, Rajan R, Sindhu RS, *et al.* Quality of life, social impact and functional outcome following ileal pouch-anal anastomosis for ulcerative colitis and familial adenomatous polyposis. *Indian J Gastroenterol* 2015;34:252–5.
 - 32 Brown C, Gibson PR, Hart A, *et al.* Long-term outcomes of colectomy surgery among patients with ulcerative colitis. *Springerplus* 2015;4:573.
 - 33 Peyrin-Biroulet L, Germain A, Patel AS, *et al.* Systematic review: outcomes and post-operative complications following colectomy for ulcerative colitis. *Aliment Pharmacol Ther* 2016;44:807–16.
 - 34 Lindsay JO, Bergman A, Patel AS, *et al.* Systematic review: the financial burden of surgical complications in patients with ulcerative colitis. *Aliment Pharmacol Ther* 2015;41:1066–78.
 - 35 Bassi A, Dodd S, Williamson P, *et al.* Cost of illness of inflammatory bowel disease in the UK: a single centre retrospective study. *Gut* 2004;53:1471–8.
 - 36 Loftus EV Jr, Friedman HS, Delgado DJ, *et al.* Colectomy subtypes, follow-up surgical procedures, postsurgical complications, and medical charges among ulcerative colitis patients with private health insurance in the United States. *Inflamm Bowel Dis* 2009;15:566–75.
 - 37 Cima RR. Timing and indications for colectomy in chronic ulcerative colitis: surgical consideration. *Dig Dis* 2010;28:501–7.
 - 38 Coward S, Heitman SJ, Clement F, *et al.* Ulcerative colitis-associated hospitalization costs: a population-based study. *Can J Gastroenterol Hepatol* 2015;29:357–62.
 - 39 Herrett E, Gallagher AM, Bhaskaran K, *et al.* Data Resource Profile: Clinical Practice Research Datalink (CPRD). *Int J Epidemiol* 2015;44:827–36.
 - 40 Wright-Hughes A, Graham E, Cottrell D, *et al.* Routine hospital data - is it good enough for trials? An example using England's Hospital Episode Statistics in the SHIFT trial of Family Therapy vs. Treatment as Usual in adolescents following self-harm. *Clin Trials* 2018;15:197–206.
 - 41 Prescription Cost Analysis – England. NHS Digital, 2017. Available: <https://digital.nhs.uk/data-and-information/publications/statistical/prescription-cost-analysis/prescription-cost-analysis-england-2017> [Accessed 1 May 2019].
 - 42 Unit costs of health and social care. Personal Social Services Research Unit, 2018. Available: <https://www.pssru.ac.uk/project-pages/unit-costs/unit-costs-2018/> [Accessed 1 May 2019].
 - 43 National tariff payment system 2017/18 and 2018/19. NHS improvement. Available: https://improvement.nhs.uk/documents/1044/2017-18_and_2018-19_National_Tariff_Payment_System.pdf [Accessed 1 May 2019].
 - 44 StataCorp. *Stata statistical software: release 15*. College Station, TX: StataCorp LLC, 2017.
 - 45 Misra R, Askari A, Faiz O, *et al.* Colectomy rates for ulcerative colitis differ between ethnic groups: results from a 15-year nationwide cohort study. *Can J Gastroenterol Hepatol* 2016;2016:8723949.
 - 46 Ahmad A, Laverty AA, Alexakis C, *et al.* Changing nationwide trends in endoscopic, medical and surgical admissions for inflammatory bowel disease: 2003-2013. *BMJ Open Gastroenterol* 2018;5:e000191.
 - 47 Kaplan GG, Seow CH, Ghosh S, *et al.* Decreasing colectomy rates for ulcerative colitis: a population-based time trend study. *Am J Gastroenterol* 2012;107:1879–87.
 - 48 Leeds IL, Truta B, Parian AM, *et al.* Early surgical intervention for acute ulcerative colitis is associated with improved postoperative outcomes. *J Gastrointest Surg* 2017;21:1675–82.
 - 49 Feuerstein JD, Curran T, Alosilla M, *et al.* Mortality is rare following elective and non-elective surgery for ulcerative colitis, but mild postoperative complications are common. *Dig Dis Sci* 2018;63:713–22.
 - 50 de Silva S, Ma C, Proulx M-C, *et al.* Postoperative complications and mortality following colectomy for ulcerative colitis. *Clin Gastroenterol Hepatol* 2011;9:972–80.