Treatment persistence and colectomy-free outcomes in patients with ulcerative colitis receiving golimumab or adalimumab: a UK experience

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ABSTRACT

Objective To examine real-world treatment persistence, colectomy-free survival and treatment switching patterns in UK patients with ulcerative colitis (UC) prescribed golimumab or adalimumab.

Design This was a retrospective chart review study in adult patients diagnosed with UC using data from 16 National Health Service sites in the UK. Patient records were included in the study if they had initiated first or second-line adalimumab or golimumab between 1 March 2016 and 30 September 2017 (index date). Subjects were required for >6 months post treatment initiation. Demographics, clinical characteristics, treatment-related data and colectomy data were extracted over a follow-up period of 6–12 months. Treatment persistence rate was the primary outcome. Colectomy-free survival and treatment switching were secondary outcomes. Outcomes were compared between treatments using χ² tests and Fisher’s exact test for categorical variables. The t-tests were used for continuous variables. Time-to-event variables were evaluated using Kaplan-Meier curves and log-rank tests.

Results The study included a total of 183 patients (96 (52.5%) prescribed adalimumab; 87 (47.5%) golimumab), and patients were mostly first-line (79.8%). Demographic and clinical characteristics were generally similar between treatment groups. Persistence rates within 12 months were 64.6% for adalimumab and 64.4% for golimumab (p=0.681). Overall, 20.2% switched to other therapy within 1 year, with 8.2% golimumab and 12.0% adalimumab switching to another biologic. Of patients prescribed adalimumab, 14.6% had ≥1 dose change, mainly dose escalations. In the 12 months post treatment initiation, 8.2% of patients underwent colectomy, with no significant difference in colectomy-free survival by treatment, p=0.73.

Conclusion This study provides evidence of clinical outcomes and real-world persistence for adalimumab and golimumab in UC. The persistence rates of both therapies were above 64.0% at 12 months following treatment initiation. In addition, the 1-year colectomy-free survival was relatively similar between the two treatments.

INTRODUCTION

Ulcerative colitis (UC) is the most common type of inflammatory bowel disease (IBD) in the UK, with around 146 000 people affected.1 UC is a chronic disease, with patients experiencing relapses and periods of remission, and patients who initially have disease of limited extent (proctitis) sometimes progressing to more extensive disease (proctosigmoiditis, left-sided colitis or pancolitis).1-3 Symptoms of UC include bloody diarrhoea, abdominal pain, urgency and tenesmus as a result of inflammation and ulceration of the colonic mucosa.2 4 5

UK and European clinical guidelines recommend that UC treatment selection should aim for sustained steroid-free remission, and consider disease severity, extent and current activity.6 7 Treatment options for UC include aminosalicylates (5-ASA), corticosteroids, immunosuppressants, biologics and the Janus kinase inhibitor, tofacitinib.8 9 Although biologics are efficacious in many patients, failure to respond has been reported to occur in 30.0%–40.0% of patients with acute refractory UC.10 A secondary loss of response to biologics is also experienced by some patients, which is believed to result from antidrug antibodies formed due to biologics being naturally produced proteins; antidrug antibodies have been detected in ≤41.0% of patients with UC receiving biologics.11

The therapeutic potential of biologics reported in clinical trials, with narrowly defined populations and carefully controlled conditions, is not always seen in clinical practice, with suboptimal adherence and persistence impacting clinical outcomes.12 Studies that explore real-world treatment outcomes following product approval are valuable in providing decision-makers with insight to the effectiveness of a product used...
What is already known about this subject?

- The therapeutic benefit observed in clinical trials for patients with ulcerative colitis (UC) treated with biologics may not necessarily be achieved in routine practice. Real-world studies are required to examine this, many of which have focused specifically on adalimumab and infliximab. US database real-world data indicates that less than half of patients continue using their initial biologic (infliximab, adalimumab, certolizumab, golimumab and vedolizumab) treatment after 1 year (44.8% in UC cohort). These persistence profiles suggest a high rate of dissatisfaction or adverse disease outcomes resulting in discontinuation and switching to a different agent. One Korean study reported at 1 and 3 years after initiation of first biologic treatment, the cumulative rates of colectomy-free survival were 100.0% and 97.3%, respectively, for adalimumab users. Another Spanish study reported probability of colectomy-free survival in golimumab patients was 85% at 52 weeks. The current study provides important data to fill the evidence gap describing key outcomes for golimumab and real-world persistence and colectomy-free data between adalimumab and golimumab in the UK.

What are the new findings?

- Analysis of data extracted from medical records for 183 patients with UC who initiated treatment with adalimumab and golimumab is reported. The persistence rates of both therapies were approximately 65.0% within 12 months following treatment initiation. Overall, 20.2% switched to other therapy within 1 year, with 8.2% golimumab and 12.0% adalimumab switching to another biologic. Colectomies were reported for 8.2% of patients in the 12 months following initiation of treatment.

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

- Awareness of the persistence levels observed in clinical practice with adalimumab and golimumab treatments for UC will support appropriate monitoring of patients and better informed decisions by clinician in the treatment of their patients, with the potential for dose escalation or treatment switching.

METHODS

Study design

This was a retrospective chart review conducted across 16 NHS sites in the UK. Each site provided data on between 2 and 30 patients, depending on the available pool of eligible patients. Data were extracted from patient medical records onto a purpose-designed standardised electronic Case Report Form (eCRF), with an eCRF completed for each eligible patient across each site. Variables included clinical characteristics, demographics, comorbidities and treatment-related data, including treatment history prior to the initiation of a biologic and details of colectomies, where present. Dose adjustment was captured but only for patients treated with adalimumab since, at the time of study design, the label for golimumab did not include dose adjustments. Subjects were required to have at least 6 months after treatment initiation up to the 12 months post initiation of the biologic if available. Overall, 86.3% (158/183) of the patients in this study had 12-month follow-up data. In order to maximise recruitment in the golimumab cohort, these patients were identified first on a consecutive basis until the site’s quota was met, or the site exhausted the pool of available patients. Following this, adalimumab patients were identified consecutively.

Patients

Inclusion criteria

According to the inclusion criteria, patients had to be at least 18 years old, and had a clinically confirmed diagnosis of UC, aligning with marketing authorisation. Patients were required to have initiated treatment with either adalimumab or golimumab at first or second line between 1 March 2016 and 30 September 2017. The date of initiation of adalimumab or golimumab was referred to as the index date. Patients were required to have data available for at least 6 months post initiation of the biologic treatment up to 12 months.

Exclusion criteria

Patients were excluded if they had received adalimumab or golimumab during the study period through a clinical trial or early access programme.

Outcomes

The primary outcome was treatment persistence rate over 6–12 months post initiation. Treatment persistence rate at any point in time was defined as the proportion of patients who had not discontinued biologic treatment at that time. The secondary outcome measures included colectomy-free survival outcome (defined as not receiving
any type of colectomy within the 6–12 months post golimumab/adalimumab treatment initiation), treatment switching pattern and dose escalation (for adalimumab).

**Statistical analysis**
Categorical variables were summarised using frequency and percentage, numeric variables using mean, SD, IQR, minimum and maximum. Variables were compared between patients receiving adalimumab and golimumab using $\chi^2$ tests for categorical variables, Fisher’s exact test for two-by-two categorical variable comparisons and t-tests for continuous variables.

Time-to-event variables were summarised using Kaplan-Meier analysis and the accompanying median (if the median was reached), and time to discontinuation and first surgery were compared between patients receiving adalimumab and those receiving golimumab. Kaplan-Meier curves were generated for treatment persistence and colectomy rates for both treatments over 12 months. Persistence and colectomy rates at 3, 6, 9 and 12 months after treatment initiation. The times at which survival outcomes was at 75.0% and 95.0% were also reported for each treatment. A log-rank test was used to determine if the persistence and colectomy rates differed statistically between treatments, with $p<0.05$ used to indicate a significant difference. To summarise treatment persistence and colectomy-free survival, Kaplan-Meier analysis was used. For persistence, the event was the treatment discontinuation and for colectomy-free survival, Kaplan-Meier analysis was used. For persistence, the event was the treatment discontinuation and for colectomy-free survival, the colectomy was the event. Time to the event was taken from the treatment initiation. Patients for which the event had not yet occurred by the date of data collection were censored at that date.

All analyses were based on observed data only, and missing data were not imputed as the number of missing values was expected to be low and there was not expected to be any systematic reason for any values that would be missing, that is, missing at random was expected. All analysis was performed using Stata Statistical software, V.15.1, College Station, Texas: StataCorp LLC.22

**RESULTS**
Patient demographic, disease and treatment characteristics are shown in table 1. Of 183 patients, 87 (47.5%) had initiated treatment with golimumab, and 96 (52.5%) with adalimumab. At least 12 months of follow-up data were available for 86.3% of the sample.

We observed a higher proportion of males treated with golimumab (71.3%) compared with adalimumab (49.0%) ($p=0.003$). Baseline characteristics between golimumab and adalimumab were relatively similar in age, body mass index (BMI), time to treatment, type of UC diseases, % treatment experienced (first line vs second line) and Charlson Comorbidity Index ($p<0.05$). At diagnosis, most patients (61.7%) had either pancolitis or left-sided colitis. The mean time between diagnosis of UC and initiation of treatment with golimumab or adalimumab were 8.5 and 7.4 years, respectively ($p=0.397$). About 80.0% of the patients received golimumab or adalimumab as their first-line biologic therapy. For 37 patients receiving a second-line biologic, half of the patients received infliximab (Remicade) as their first-line treatment followed by infliximab (biosimilar) (37.8%). Antidrug antibody status after first-line treatment was available for very few patients receiving a second-line biologic. Overall, 24.3% of patients tested positive for antidrug antibodies at discontinuation of their first-line biologic treatment.

**Treatment persistence**
At 12 months following treatment initiation, 56 (64.4%) patients receiving golimumab and 62 (64.6%) patients receiving adalimumab remained on treatment. Kaplan-Meier analysis revealed that there was no significant difference in persistence rates over 12 months between golimumab and adalimumab ($p=0.681$; figure 1). However, we found that proportions of patients receiving golimumab remained on treatment at 3, 6, 9 and 12 months after treatment initiation were slightly higher, although not statistically significant, than those receiving adalimumab (figure 1 and table 2). The time for medication persistence rates at 75.0% was slightly longer for golimumab (8.6 months) compared with adalimumab (6.8 months) (table 2). The sensitivity analysis adjusting for sex revealed no significant difference of treatment persistence between golimumab and adalimumab patients.

Regarding the data on antidrug antibodies, only 17 (26.2%) patients who discontinued treatment had the results of tests for antidrug antibodies recorded (6 receiving golimumab and 11 receiving adalimumab); a positive result was reported for only one patient, who was receiving adalimumab.

**Treatment switching**
Overall, 20.2% (n=37) switched to other therapy within 1 year. By treatment, 8.2% (n=15) of patients receiving golimumab switched to another biologic compared with 12.0% (n=22) of those receiving adalimumab. Among patients switching biologics (n=37), 56.8% switched to vedolizumab, followed by infliximab biosimilar (29.7%) and infliximab bio-originator (13.5%) (table 2). The switching pattern for both medications was similar. Of 22 patients initially prescribed adalimumab who switched to another biologic, 14 (63.6%) switched to vedolizumab, 6 (27.3%) to an infliximab biosimilar and 2 (9.1%) to infliximab bio-originator. Of those 15 patients initially prescribed golimumab, more than half switched to either infliximab biosimilar or to infliximab bio-originator (table 2).

Data on dose adjustment were available for 90 patients prescribed adalimumab (Table not shown). We found that 14 (14.6%) had at least one dose adjustment. The majority (85.7%) of cases had a dose escalation. Kaplan-Meier analysis indicated that 10.0% of adalimumab
Table 1  Patient demographic, clinical and treatment characteristics

|                             | Total   | Adalimumab | Golimumab | P value |
|-----------------------------|---------|------------|-----------|---------|
|                             | N=183   | N=96       | N=87      |         |
| Age at treatment initiation (years) |         |            |           |         |
| N                           | 183     | 96         | 87        | 0.2971  |
| Mean (SD)                   | 45.6 (15.0) | 44.4 (15.6) | 46.8 (14.3) |         |
| Median                      | 45.3    | 42.7       | 47.1      |         |
| Sex, n (%)                  |         |            |           | 0.0026  |
| N                           | 183     | 96         | 87        |         |
| Female                      | 74 (40.4) | 49 (51.0)  | 25 (28.7) |         |
| Male                        | 109 (59.6) | 47 (49.0)  | 62 (71.3) |         |
| BMI (kg/m²)                 |         |            |           | 0.7936  |
| N                           | 55      | 17         | 38        |         |
| Mean (SD)                   | 26.95 (5.79) | 26.64 (5.95) | 27.09 (5.79) |         |
| Median                      | 25.83   | 25.51      | 26.41     |         |
| Charlson Comorbidity Index  |         |            |           | 0.0503  |
| N                           | 183     | 96         | 87        |         |
| Mean (SD)                   | 0.10 (0.42) | 0.16 (0.55) | 0.03 (0.18) |         |
| Median                      | 0.00    | 0.00       | 0.00      |         |
| Disease extent at diagnosis, n (%) |         |            |           | 0.3384  |
| N                           | 183     | 96         | 87        |         |
| Ulcerative proctitis        | 26 (14.2) | 11 (11.5)  | 15 (17.2) |         |
| Proctosigmoiditis           | 12 (6.6) | 9 (9.4)    | 3 (3.4)   |         |
| Left-sided colitis          | 54 (29.5) | 26 (27.1)  | 28 (32.2) |         |
| Pancolitis                  | 59 (32.2) | 31 (32.3)  | 28 (32.2) |         |
| Do not know                 | 32 (17.5) | 19 (19.8)  | 13 (14.9) |         |
| Time between UC diagnosis and treatment initiation (months) |         |            |           | 0.3974  |
| N                           | 168     | 84         | 84        |         |
| Mean (SD)                   | 95.4 (107.6) | 88.3 (114.0) | 102.4 (101.0) |         |
| Median                      | 58.1    | 43.9       | 63.7      |         |
| Treatment line, n (%)       |         |            |           | 0.7129  |
| N                           | 183     | 96         | 87        |         |
| First-line biologic         | 146 (79.8) | 78 (81.3)  | 68 (78.2) |         |
| Second-line biologic        | 37 (20.2) | 18 (18.8)  | 19 (21.8) |         |
| First-line biologic for patients at second line, n (%) |         |            |           | 0.4022  |
| N                           | 37      | 18         | 19        |         |
| Adalimumab                  | 1 (2.7) | 0 (0.0)    | 1 (5.3)   |         |
| Golimumab                   | 1 (2.7) | 1 (5.6)    | 0 (0.0)   |         |
| Infliximab (bio-originator) | 19 (51.4) | 10 (55.6)  | 9 (47.4)  |         |
| Infliximab (biosimilar)     | 14 (37.8) | 7 (38.9)   | 7 (36.8)  |         |
| Other                       | 2 (5.4) | 0 (0.0)    | 2 (10.5)  |         |
| Antidrug antibodies after first-line biologic, n (%) |         |            |           | 0.4501  |
| N                           | 37      | 18         | 19        |         |
| Positive                    | 9 (24.3) | 6 (33.3)   | 3 (15.8)  |         |
| Negative                    | 10 (27.0) | 4 (22.2)   | 6 (31.6)  |         |
| Not known/not tested        | 18 (48.6) | 8 (44.4)   | 10 (52.6) |         |

BMI, body mass index; IQR, inter-quartile range; SD, standard deviation; UC, ulcerative colitis.
patients had undergone a first dose adjustment at 7.3 months.

Colectomy-free outcomes
A total of 8.2% (n=15) of patients underwent colectomy within 12 months following initiation of golimumab or adalimumab. At 12 months, 88 (91.7%) and 80 (92.0%) patients who received adalimumab and golimumab, respectively, demonstrated colectomy-free survival at 12 months. The Kaplan-Meier analysis showed no significant difference between the two treatments in colectomy-free survival over the 12 months after treatment initiation (p=0.735; figure 2). Our sensitivity analysis when adjusting for sex confirmed the colectomy-free survival result. When evaluating the proportion of patients at 3,6,9 and 12 months, patients with golimumab were slightly higher, although not statistically significant, than those receiving adalimumab at 12 months (94.2% vs 91.7%, respectively) after treatment initiation (table 3). In addition, the average time for colectomy-free survival rates to drop to 95.0% was slightly longer for golimumab (8.5 months) compared with adalimumab (6.1 months); however, this was not statistically significant (table 3).

Table 2  Kaplan-Meier analysis of persistence and treatment switching data

|                          | Total     | Adalimumab | Golimumab |
|--------------------------|-----------|------------|-----------|
| Persistence data         |           |            |           |
| Missing, n               | 5         | 6          |           |
| Events, n                | 30        | 25         |           |
| Censored, n              | 61        | 56         |           |
| Time from treatment initiation (months) |     |            |           |
| 3                        | 85.7      | 90.1       |           |
| 6                        | 76.9      | 80.2       |           |
| 9                        | 69.2      | 72.8       |           |
| 12                       | 67.0      | 69.1       |           |
| 75% of patients remaining on treatment | 6.8       | 8.6        |           |
| Switching data           |           |            |           |
| Per cent of patients who stopped adalimumab or golimumab within 12 months, n (%) | 65 (35.5) | 34 (35.4) | 31 (35.6) |
| Per cent of patients who switched to another biologic within 12 months, n (%) | 37 (20.2) | 22 (12.0) | 15 (8.2) |
| Of those who switched, which biologic did they switch to? n (%) |            |            |           |
| N                        | 37        | 22         | 15        |
| Infliximab (Remicade)    | 5 (13.5)  | 2 (9.1)    | 3 (20.0)  |
| Infliximab biosimilar (Inflectra/Remsima) | 11 (29.7) | 6 (27.3)  | 5 (33.3)  |
| Vedolizumab (Entyvio)    | 21 (56.8) | 14 (63.6)  | 7 (46.7)  |
A substantial proportion of patients used additional therapies for their UC in addition to a biologic during the 12 months study period (data not shown). Oral 5-ASA was the most commonly prescribed therapy, taken by 41.4% and 46.9% of patients receiving golimumab and adalimumab, respectively. Corticosteroids were prescribed to 35.6% (golimumab) and 39.6% (adalimumab), respectively, and immunomodulators to 26.4% and 25.0%, respectively. Only 50 (27.3%) patients overall had no additional therapy reported in the 12 months post biologic initiation, with 23 (26.4%) patients prescribed golimumab and 27 (28.1%) patients being prescribed adalimumab. The proportions of patients receiving additional (or no) UC medications were similar between patients receiving golimumab and adalimumab.

**DISCUSSION**

This study was a retrospective chart review analysis of 183 UK patients. The persistence rate was approximately 65.0% at 12 months post treatment initiation in patients prescribed the anti-TNF agents golimumab and adalimumab, respectively. In the real world, persistence may be viewed as a surrogate measure of drug efficacy. For golimumab, in the phase 3 Program of Ulcerative Colitis Research Studies Utilizing an Investigational Treatment (PURSUIT) trial, clinical response was maintained through week 54 in approximately 50.0% of patients, all of whom were anti-TNF naïve.23 Sandborn et al conducted a subgroup analyses on the Ulcerative colitis long-term remission and maintenance with adalimumab 2 (ULTRA 2) clinical trial data to evaluate the 1-year maintenance outcomes among patients with moderately-to-severely active UC who responded to induction therapy with adalimumab. The study showed that 30.9% of patients achieved clinical remission and 49.6% achieved clinical response at week 52 (12 months).24 A real-world postmarketing study in the UK showed that, of 205 anti-TNF naïve patients receiving golimumab, 68.8% achieved clinical response rate at week 6% and 38.5% had clinical remission.25 Our study showed that the persistence rates of the two medications slightly higher than what we observed in clinical trials. These real-world persistence results provide a broader perspective that can be used to aid treatment decisions in a more heterogenous clinical setting. Our analysis demonstrated that discontinuation of treatment did not appear to result from the known development of antidrug antibodies, as testing was reported in about a quarter of patients, with a positive test reported for only one patient receiving adalimumab.

Real-world studies on treatment persistence reported from other countries ranged from 35.0% to 85.0%.13 14 17 26 27 A recent real-world study from Canada reported 63.0% of patients persisted with golimumab treatment.17 A retrospective study using US claims data reported overall persistence rates of 59.0% at 1 year in biologic-naïve patients with UC, with 56.0% and 44.0% of patients prescribed adalimumab and golimumab, respectively.16 A real-world analysis from a large-scale US database reported that of patients newly diagnosed with UC prescribed biologic treatment (ie, adalimumab, certolizumab, golimumab, infliximab or vedolizumab), 45.0% persisted with treatment 1 year after initiation.13 One-year persistence rates in this US study for adalimumab and golimumab were 45.0% and 40.0%, respectively. Overall, 54.0% of patients with UC newly initiated on a biologic (adalimumab, certolizumab, golimumab, infliximab, natalizumab, ustekinumab or vedolizumab) were reported to remain on their first-line therapy at 1 year based on another analysis of the same US insurance claims database.15 However, McDermott et al reported only 35.0% (n=8/23) of patients receiving adalimumab persisting at 1 year.14 This Irish study might be linked to the low patient numbers, as only 23 out of 3000 patients were found to be patients with UC receiving adalimumab with a high percentage of previous biologic failure (20 of 23 patients had previously received infliximab).

### Table 3 Colectomy-free survival analysis

|                | Adalimumab | Golimumab |
|----------------|------------|-----------|
| N              | 96         | 87        |
| Missing, n     | 0          | 1         |
| Events, n      | 8          | 6         |
| Censored, n    | 88         | 80        |
| Time from treatment initiation (months) | Patients remaining colectomy free, (%) |
| 3              | 99.0       | 97.7      |
| 6              | 95.8       | 96.5      |
| 9              | 94.8       | 94.2      |
| 12             | 91.7       | 94.2      |
| 95% of patients remaining colectomy free | Time (months) |
|                | 6.1        | 8.5       |
The literature from a recent review of surgery in patients with IBD reported that, although biologics may delay the need for colectomy, 10.0%–30.0% of patients with UC will ultimately require surgery. Our analysis of patient records showed that about 8.0% of patients requiring colectomy in the 12 months following the initiation of golimumab and adalimumab. Other studies reported in the literature support high rates of colectomy-free survival in patients receiving these biologics. Given that data were only captured for a maximum of 12 months, in the present study, it is unknown what proportion of these patients may go on to require colectomy at a later timepoint. Two studies in Italian primary IBD centres reported short-term (3 months) colectomy rates of 1.0% and 3.0% in patients with UC receiving golimumab and adalimumab, respectively. Two retrospective Spanish multicentre cohort studies reported colectomy outcomes in approximately 16 (11.0%) of 142 patients with UC receiving golimumab, compared with approximately 22 (12.0%) of 184 patients with adalimumab as maintenance therapy, during a median 23-month follow-up period and in our sensitivity analysis when adjusting for sex, revealed no significant difference of treatment persistence or colectomy-free survival between golimumab and adalimumab patients.

A systematic literature review reports that switching rate in anti-TNF ranged from approximately 4 (1.0%) of 380 patients at 6 months to approximately 140 (26.0%) of 538 patients at 2 years. In addition, they reported the most common switching pattern, infliximab to adalimumab, occurred in 3.8% (median 5.6 years) to 25.5% (mean 3.3 years) of patients. A US study reported switch rate to another biologic ranged from 4.5% to 20%. About 20.0% of patients switched to other biologic in our study. The most common biologic during the study period that patients switched to was vedolizumab, followed by infliximab.

In our study, 15.0% of patients receiving adalimumab required a dose adjustment within 1 year, and the majority of adjustments were dose escalation. The literature reports a wide range of rates of dose escalation for biologics in UC. Dose escalation of adalimumab patients ranged from 16.0% to 43.0% of patients at 1 year. A Canadian retrospective study of 136 patients with UC receiving golimumab maintenance therapy reported 5.0% having their dose escalated, while an observational study conducted in 14 Italian primary IBD centres reported dose escalations in approximately 17 (16.0%) of 107 patients with UC receiving adalimumab, similar to our findings. The lower rates of dose escalation for golimumab may be due to the lack of approval for dose adjustments for this drug. An analysis of NHS Hospital Episode Statistics in England reported approximately 82 (43.0%) of 191 patients with UC required adalimumab dose escalation during the maintenance phase; however, the duration during which patient data were observed was not reported. Due to the paucity of data, the long-term outcome of dose escalation needs further studies.

To our knowledge, this is the first study comparing persistence and colectomy outcomes between golimumab and adalimumab treatment in patients with UC in the UK. In this cohort, patient demographics such as age, BMI, time to treatment and type of UC diseases as well as their comorbidity profile were comparable between the two treatment groups, with the exception of gender (higher proportion of males in the golimumab cohort). Although treatment was not randomised in this observational study, the impact of potential confounding factors could be low due to the small sample size. The sample size precluded carrying out adequate subgroup analysis between patients receiving golimumab or adalimumab as first-line or second-line anti-TNF. However, the results of our study must be interpreted taking into consideration the known limitations of retrospective real-world data studies. In particular, some degree of missing data occurred due to incomplete medical records, assumed subtotal colectomy as an exclusion criterion and also end point of colectomy-free survival, and all analyses were based on observed data only. Therefore, the number of patients for each variable was varied. Finally, at the time this study was conducted, dose escalations for golimumab were not approved. However, currently early dose escalation of golimumab is approved by the European Medicines Agency, and real-world data are now available, reporting that early dose optimisation of golimumab improves clinical outcomes. If golimumab dose escalation had been possible to include in this study, clinical outcomes observed may have been better. Despite these limitations, we believe that these real-world data, obtained from 16 sites in the UK to avoid single centre bias, provide a useful insight to treatment persistence and clinical outcomes in patients with UC prescribed the anti-TNF agents golimumab and adalimumab.

CONCLUSIONS

Real-world data on treatment persistence and clinical outcomes in clinical settings are essential to understand the real-world effectiveness of pharmaceutical interventions. Our study evaluated important data to fill the evidence gap comparing real-world persistence and colectomy-free outcomes for adalimumab and golimumab in the UK. The persistence rates of both therapies were above 64.0% at 12 months following treatment initiation. In addition, the 1-year colectomy-free survival was relatively similar between the two treatments. Additional research to evaluate long-term persistence and clinical outcomes in the UK should be encouraged.

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Patient consent for publication Not required.

Ethics approval The study was conducted in accordance with Good Pharmacoepidemiology Practices issued by the International Society for Pharmacoepidemiology. The study documentation was submitted and approved by the Health Research Authority and a UK National Health Service Research Ethics Committee.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request. Data collection was undertaken by Adelphi Real World as part of a real-world non-interventional retrospective chart review study, entitled ‘Treatment persistence and colonic remission-free outcomes in patients with ulcerative colitis receiving golimumab or adalimumab: A UK experience’, sponsored by Merck & Co. All data that support the findings of this study are the intellectual property of Merck & Co. All requests for access should be addressed directly to Merck & Co. AP at amy@merck.com.

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REFERENCES
1. (NICE). N.I.F.H.a.C.E. Inflammatory bowel disease, 2015. Available: https://www.nice.org.uk/guidance/gs81/chapter/Introduction
2. Ungaro R, Mehandru S, Allen PB, et al. Ulcerative colitis. Lancet 2017;389:1756–70.
3. Ondás I, Eckmann L, Talamini M, et al. Ulcerative colitis. Lancet 2012;380:1606–19.
4. Antonelli E, Villanacci V, Bassotti G. Novel oral-targeted therapies for mucosal healing in ulcerative colitis. World J Gastroenterol 2018;24:5322–30.
5. Feuerstein JD, Cheifetz AS. Ulcerative colitis: epidemiology, diagnosis, and management. Mayo Clin Proc 2014;89:1553–63.
6. (NICE). N.I.F.H.a.C.E. Interactive flowchart: inducing remission in people with ulcerative colitis overview, 2019. Available: https://pathways.nice.org.uk/pathways/ulcerative-colitis#path-way%3A/pathways/ulcerative-colitis/inducing-remission-in-people-with-ulcerative-colitis.xml!content=view-index [Accessed 18 Dec 2019].
7. Harbord M, Eliakim R, Bettenworth D, 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.
8. Danese S, Aliez M, van Bodegraven AA, et al. Unmet medical needs in ulcerative colitis: an expert group consensus. Dig Dis 2019;37:266–83.
9. Wiekamp J, Stange EF. Recent advances and emerging therapies in the non-surgical management of ulcerative colitis. F1000Res 2018;7.
10. Lim AW, Panaccione R, Seow CH. Exploring the role of monitoring anti-TNFα drug and antibody levels in the management of inflammatory bowel disease. Therap Adv Gastroenterol 2011;4:145–61.
11. Vermeire S, Gils A, Accossato P, et al. Immunogenetics of biologics. Therap Adv Gastroenterol 2018;11:1758283x17750355.
12. Maniadiakis N, Toth E, Schiff M, et al. A targeted literature review examining biologic therapy compliance and persistence in chronic inflammatory diseases to identify the associated unmet needs, driving factors, and consequences. Adv Ther 2018;35:1333–55.
13. Chen C, Hartzema AG, Xiao H, et al. Real-World pattern of biologic use in patients with inflammatory bowel disease: treatment persistence, switching, and importance of concurrent immunosuppressive therapy. Inflamm Bowel Dis 2019;25:1417–27.
14. McDermott E, Murphy S, Keegan D, et al. Efficacy of adalimumab as a long-term maintenance therapy in ulcerative colitis. J Crohns Colitis 2015;7:150–63.
15. Brady JE, Stott-Miller M, Mu G, et al. Treatment patterns and sequencing in patients with inflammatory bowel disease. Clin Ther 2018;40:1509–21.
16. Comerford E DM, Smith T, et al. Treatment patterns of biologic therapies used to treat ulcerative colitis: a retrospective database analysis in the United States. Am J Gastroenterol 2019;114:pS7.
17. Bressler B, Williamson M, Sattin B, et al. Real world effectiveness of golimumab therapy in ulcerative colitis regardless of prior TNF exposure. J Can Assoc Gastroenterol 2018;1:129–34.
18. Taxonera C, Rodriguez C, Bertoletti F, et al. Clinical outcomes of golimumab as first, second or third anti-TNF agent in patients with moderate-to-severe ulcerative colitis. Inflamm Bowel Dis 2017;23:1394–402.
19. Taxonera C, Iglesias E, Muñoz F, et al. Adalimumab maintenance treatment in ulcerative colitis: outcomes by prior anti-TNF use and efficacy of dose escalation. Dig Dis Sci 2017;62:481–90.
20. Tursi A, Allegretta L, Buccianti N, et al. Effectiveness and safety of golimumab in treating outpatient ulcerative colitis: a real-life prospective, multicentre, observational study in primary inflammatory bowel diseases centers. J Gastroenterol Hepatol 2017;32:239–44.
21. Otsuka T, Ooi M, Tobimatsu K, et al. Short-term and long-term outcomes of infliximab and tacrolimus treatment for moderate to severe ulcerative colitis: retrospective observational study. Kobe J Med Sci 2018;64:E140–8.
22. StataCorp. Stata Statistical Software: Release 15. College Station, TX: StataCorp LLC, 2017.
23. Sandborn WJ, Feagan BG, Marano C, et al. Subcutaneous golimumab maintains clinical response in patients with moderate-to-severe ulcerative colitis. Gastroenterology 2014;146:96–109.
24. Sandborn WJ, Colombeau J-F, D’Haens G, et al. One-Year maintenance outcomes among patients with moderately-to-severely active ulcerative colitis who responded to induction therapy with adalimumab: subgroup analyses from ultra 2. Aliment Pharmacol Ther 2013;37:204–13.
25. DOP049. Efficacy and safety of golimumab induction for moderate-to-severe ulcerative colitis in the United Kingdom: results from the GO-COLITIS study. Journal of Crohn’s and Colitis 2016;10:S57–8.
26. Angelson L, Almer S, Davidstott L, et al. Short and long-term efficacy of adalimumab in ulcerative colitis: a real-life study. Scand J Gastroenterol 2020;55:141–62.
27. Patel H, Latremouille-Viau D, Burne R, et al. Comparison of real-world treatment outcomes with Vedolizumab versus infliximab in Biologic-Naive patients with inflammatory bowel disease. Crohns Colitis 360 2019;1:360.
28. Wong DJ, Roth EM, Feuerstein JD, et al. Surgery in the age of biologics. Gastroenterol Rep 2019;7:77–90.
29. Armuzzi A, Pugliese D, Nardone OM, et al. Management of difficult-to-treat patients with ulcerative colitis: focus on adalimumab. Drug Des Devel Ther 2013;7:289–96.
30. Tursi A, Eiesser B, Mohan W, et al. Effectiveness and safety of adalimumab to treat outpatient ulcerative colitis: a real-life multicenter, observational study in primary inflammatory bowel disease centers. Medicine 2018;97:e11897.
31. Gemayel NC, Rizzello E, Atanasov P, et al. Dose escalation and switching of biologics in ulcerative colitis: a systematic literature review in real-world evidence. Curr Med Res Opin 2019;35:1911–23.
32 Khan S, Rupniewska E, Neighbors M, et al. Real-World evidence on adherence, persistence, switching and dose escalation with biologics in adult inflammatory bowel disease in the United States: a systematic review. *J Clin Pharm Ther* 2019;44:495–507.

33 Black CM, Yu E, McCann E, et al. Dose escalation and healthcare resource use among ulcerative colitis patients treated with adalimumab in English hospitals: an analysis of real-world data. *PLoS One* 2016;11:e0149692.

34 Taxonera C, Iborra M, Bosca-Watts MM, et al. Early dose optimization of golimumab induces late response and long-term clinical benefit in moderately to severely active ulcerative colitis. *Curr Med Res Opin* 2019;35:1297–304.