

Impact of Integrated
Health System
Specialty Pharmacy
Services and
Associated Patient
Factors on
Inflammatory Bowel
Disease Outcomes

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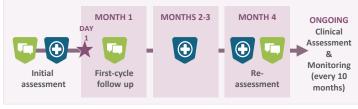
#### DISCLOSURES

The authors of this presentation have nothing to disclose concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the white matter of this presentation.

### BACKGROUND

- Inflammatory bowel disease (IBD) encompasses Crohn's disease (CD) and ulcerative colitis (UC). While CD and UC are clinically diverse, both are characterized by chronic inflammation of the gastrointestinal tract.
- Research suggests that a treat-to-target approach be used to manage this
  population, utilizing subjective and objective markers of disease.<sup>1</sup>
- Literature exists to support the role of a clinical pharmacy team in improving medication access, adherence, and quality of care; however, there are limited data on the pharmacy role in improving IBD outcomes.<sup>2-4</sup>
- This study aims to assess the impact of integrated health system specialty pharmacy [HSSP] services on IBD outcomes.

#### Figure 1: HSSP IBD Patient Journey









Care Liaison Interaction

### **METHODS**

Study Design: Multi-center, retrospective observational analysis of adult and pediatric IBD patients receiving biologic or small molecule agents from HSSPs between January 1, 2022 and December 31, 2023

Inclusion Criteria: Patients enrolled in the HSSP services for ≥ 4 months with a baseline and follow-up assessment of corticosteroid use, flares, and pain scores. Patients with ICD-10 codes unrelated to CD and UC were excluded.



**Primary Outcome:** percent reduction in corticosteroid usage from baseline

Secondary Outcome: percent reduction in IBD flares from baseline and reduction in average pain score from baseline



**Data Identification:** Data collected included age, sex, IBD medication, ICD-10 code, primary insurance type, treatment status, out-of-pocket cost, days on service, medication adherence measured by the proportion of days covered (PDC), corticosteroid use, number of IBD flares, and pain severity.



Analysis: A logistic regression model was utilized to evaluate the impact of various factors on changes in steroid use, flares, and pain.

REFERENCES

1. Turner D, Riciculto A, Levis A, et al. STRIDE-II: An Update on the Selecting Therapeutic Targets in Inflammatory Bowel Disease (STRIDE) initiative of the International Organization for the Study of IBD (1088): Determining Therapeutic Goals for Treat-to-Target strategies in IBD. Gastroenterology. 2021;16(5)(5)(1570-1583. doi:https://doi.org/10.1053/j.gastro.2020.12.011
2. CHO LOR Bullo P. Paraparamola IA. And. Robe entire Insert of a Clinical Pharmace Rowel Disease Center. Cordor's & Gallot 2023-57(1)(40810).

. Alrashed F, Almutairi N, Shehab M. The Role of Clinical Pharmacists in Improving Quality of Care in Patients with Inflammatory Bowel Disease: An Evaluation of Patients' and Physiciatisfaction. Healthcare. 2022;10(10):1818. doi:https://doi.org/10.3390/healthcare10101818

Danielle Mae Thank, Kandilian R, Khanh Le Nguyen, P099 Implementation of an Inflammatory Bowel Disease Clinical Pharmacy Service. The American Journal of Gastroenterology 19;114(1):526-526. doi:https://doi.org/10.14309/01.ajg.0000613364.78413.id.

### **RESULTS**

**Table 1** summarizes patient characteristics and associated patient factors influence on steroid use, flares, and pain. The regression model showed that steroid-free and symptom-free patients at baseline were more likely to maintain positive outcomes at follow-up. We observed a 69% reduction in corticosteroid (**Figure 2**), 62% decrease in disease flares (**Figure 3**), and an average decrease in pain scores of 16% (**Figure 4**). The mean PDC was 94.5%.

Table 1: Patient Characteristics and Associated Patient Factors

Characteristic	N = 1373	Steroid Use	Flares	Pain
Age (n, %) <65 ≥65	1198 (87%) 175 (13%)	0.123	0.150	0.632**
Sex (n, %) M F Unknown	655 (48%) 688 (50%) 30 (2%)	<b>0.361*</b> - -0.448	<b>0.422***</b> - 0.439	0.123 - 0.713
Diagnosis (n, %) UC CD	398 (29%) 975 (71%)	-0.627*** -	-0.180	0.381**
Treatment Status (n, %) Experienced Naïve	1119 (82%) 254 (18%)	0.173	- 0.071	0.037
Steroid Free at Baseline (n, %)	1061 (77%)	2.115***		
Symptom Free at Baseline (n, %)	882 (64%)		0.940***	1.845***
Days of Service <sup>2</sup> (Range)	428 (129-727)	-0.002**	0.001*	0.0002
Insurance Type (n, %) Commercial Medicaid Medicare Unknown/Other	629 (46%) 105 (8%) 209 (15%) 430 (31%)	- -0.558 -0.162 0.345	-0.272 -0.103 0.091	- -0.255 -0.255 0.057
Copay (n, %) > \$0 = \$0  'n<0.1: 'n<0.05: '''n<0.01: 2 Median	527 (38%) 846 (62%)	0.143	-0.118	0.138

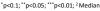


Figure 2: Corticosteroid Use

400
312
300
200
95
81
73
Total
Population Naïve Experience

Baseline

■ Follow Up

Figure 3: IBD Flares

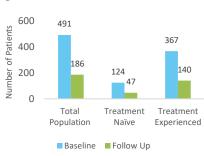
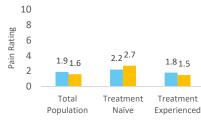


Figure 4: Average Pain



# ■ Baseline ■ Follow Up

## **CONCLUSIONS**

- IBD patients enrolled in HSSP services demonstrated clinically meaningful responses illustrated by the reduction in corticosteroid use, IBD flares, and average pain score.
- Patients achieved a consistently high adherence rate of 94.5%. This high adherence rate suggests the value of the HSSP in promoting adherence to specialty therapies.
- Steroid-free and symptom-free status at baseline are associated with positive outcomes, but additional analysis is needed to better identify what factors contribute to IBD outcomes.
- These findings highlight the potential for sustained disease control and improved quality of life. Additionally, they
  contribute to mitigating the risks associated with long-term corticosteroid use.