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#### Background

- Inflammatory bowel disease (IBD) is associated with substantial economic burden and its increasing prevalence will only exacerbate the costs.<sup>1,2</sup>
- Population health management (PHM) strategies like risk stratification are needed to increase quality of care and improve health outcomes while reducing costs.<sup>3</sup>
- Risk stratification is also integral for IBD patient management according to care pathways/guidelines.<sup>4-8</sup>
- Previous research suggests risk stratification using American Gastroenterological Association (AGA) care pathways was infrequently documented.
- IBD PATH was developed to identify patient risk and potential gaps in care using electronic medical record (EMR) data.
- We used IBD PATH to conduct a real-world case study to identify data standardization gaps and facilitate PHM efforts within Ochsner Health (Ochsner).

### Materials and Methods

- Ochsner data included patients with IBD visits between Jan. 2020 and Dec. 2021 and data variables; medication name and ordered date, visit dates and associated diagnosis and procedure codes.
- Using a standardized template included in IBD PATH, a subset of patients had additional unstructured clinical data from EMR converted into structured data.
- These variables are based on AGA care pathway risk factors for Crohn's disease (CD) and ulcerative colitis (UC) complications.<sup>4,5</sup>
- The EMR dataset, formatted per the tool specifications, was uploaded into the tool.
- Descriptive analyses were performed through IBD PATH.

#### Results

- AGA risk was not documented in the EMR. Unstructured risk factors were collected for a total of 164 patients (124 CD, 71 UC), Table 1.
- Majority of cases (82% CD, 77% UC) were classified as mod/high risk and the primary risk factors perianal disease for CD and previous steroid-requiring disease for UC.

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# Inflammatory Bowel Disease Population Analyzer Tool for Health systems (IBD PATH): A Case Study Risk Stratifying IBD Patients for Clinical Outcomes

### Table 1: Respondent Demographics

-	•							
		Crohn's Disease		Ulcerative Colitis				
	Total, N=124	Mod/High Risk, N=102	Low Risk, N=22	Total, N=71	Mod/High Risk, N=55	Low Risk, N=16		
AGA Risk Level Characteristics, n(%)								
Crohn's Disease Risk Factors								
Age <30 years at diagnosis	22 (18)	22 (22)						
Extensive anatomic involvement	7 (6)	7 (7)						
Perianal disease	83 (67)	83 (81)						
Severe rectal disease	32 (26)	32 (31)						
Deep ulcers	9 (7)	9 (9)						
Previous surgical resection	0 (0)	0 (0)						
Stricturing behavior	17 (14)	17 (17)						
Penetrating behavior	24 (19)	24 (24)						
Ulcerative Colitis Disease Risk Factors								
Age <40 years				16 (23)	16 (29)			
Extensive colitis				33 (46)	33 (60)			
Steroid-requiring disease				43 (61)	43 (78)			
Deep ulcers				2 (3)	2 (4)			
History of hospitalization				4 (6)	4 (7)			
High CRP and ESR				0 (0)	0 (0)			
Clostridium difficile infection				4 (6)	4 (7)			
Cytomegalovirus infection				1 (1)	1 (2)			
Medication Utilization, n(%)								
Biologics	38 (31)	35 (92)	3 (8)	16 (23)	16 (100)	7 (13)		
Adalimumab	12 (32)	10 (83)	2 (17)	5 (31)	5 (100)	0 (0)		
Certolizumab pegol	0 (0)	0 (0)	0 (0)	1 (6)	1 (100)	0 (0)		
Golimumab				1 (6)	1 (100)	0 (0)		
Infliximab	14 (56)	14 (100)	0 (0)	2 (13)	2 (100)	0 (0)		
Vedolizumab	10 (26)	9 (90)	1 (10)	5 (31)	5 (100)	0 (0)		
Ustekinumab	3 (8)	3 (100)	0 (0)	3 (19)	3 (100)	0 (0)		
Immunomodulators	5 (4)	3 (60)	2 (40)	4 (6)	3 (75)	1 (25)		
Azathioprine	5 (100)	3 (60)	2 (40)	4 (100)	3 (75)	1 (25)		
5-Aminosalicylic acids				12 (17)	9 (75)	3 (25)		
Mesalamine				10 (83)	7 (70)	3 (30)		
Sulfasalazine				2 (17)	2 (100)	0 (0)		

**Note:** The medication utilization percentages for the mod/high and low risk columns are based on total number of patients in the row not total number of mod/high or low risk patients



Note: The quality indictors percentages for the mod/high and low risk columns are based on total number of patients in the row not total number of mod/high or low risk patients Abbreviations: Mod, moderate; AGA, American Gastroenterological Association; CRP, c-reactive protein; ESR, erythrocyte sedimentation rate

#### Contact

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### References

- Dis. 2020:26(4):619-625.
- 3. Girwar SM, et al. A systematic review of risk stratification tools internationally used in primary care settings. *Health Sci Rep.* 2021;4(3):e329.
- 4. Sandborn WJ. Crohn's disease evaluation and treatment: Clinical decision tool. *Gastroenterology*. 2014;147(3):702-703.
- 5. Dassopoulos T, et al. Ulcerative colitis care pathway. *Gastroenterology*. 2015;149(1):238-245.
- 6. Lichtenstein GR, et al. ACG clinical guideline: management of Crohn's disease in adults. Am J Gastroenterol. 2018;113(4):481-517.
- 7. Rubin DT, et al. ACG clinical guideline: ulcerative colitis in adults. Am J Gastroenterol. 2019;114(3):384-413.
- 9. Alli-Akintade L, Pruthvi P, Hadi N, Sachar D. Race and fistulizing perianal crohn's disease. Journal of *Clinical Gastroenterology*. 2015;49(3):e21-e23.

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- While 30% of mod/high risk patients had a biologic medication record, nearly 50% did not have an IBD treatment record.
- Mod/high risk patients were also more likely to have had IBD-related hospitalizations, emergency department visits, and procedures in the follow-up period.
- Steroid use was low overall; however, majority of those that had evidence of steroid use were mod/high risk. Additionally, narcotic analgesic use was high in those with mod/high risk.

## Figure 1: Patient-Level Report

PATIENT LEVEL REPORT											
Type Any	•	Risk Level Any 🔻		Medications Any		<ul> <li>Apply Filter</li> </ul>		View Excel Report			
Patient	Provider	IBD Type	Risk Level	User Defined 1	User Defined 2	User Defined 3	Aminosalicylates	Biologics	JAK inhibitor	Immunomodulators	Steroids
291	1234	CD	Mod/High				No	No	No	No	No
296	1234	UC	Low				No	No	No	No	No
32	1234	UC	Mod/High				No	No	No	No	No
332	1234	Both	Mod/High				No	No	No	Azathioprine	No
339	1234	Both	Mod/High				No	Adalimumab	No	No	No
347	1234	UC	Mod/High				No	No	No	No	Prednisone
352	1234	Both	Mod/High				No	Adalimumab	No	No	No
361	1234	Both	Mod/High				No	Infliximab	No	No	No
367	1234	CD	Mod/High				No	No	No	No	No
378	1234	UC	Low				No	No	No	No	No
413	1234	Both	Mod/High				No	Infliximab	No	No	No
414	1234	CD	Mod/High				No	Infliximab	No	No	No
420	1234	CD	Mod/High				No	No	No	No	No
423	1234	Both	Mod/High				Mesalamine	No	No	Azathioprine	No
425	1234	UC	Low				No	No	No	No	No
440	1234	UC	Mod/High				No	No	No	No	No
475	1234	CD	Mod/High				No	Infliximab	No	No	No
486	1234	CD	Mod/High				No	No	No	No	No
509	1234	UC	Mod/High				No	Ustekinumab	No	No	No
514	1234	Both	Mod/High				No	No	No	No	Prednisone
529	1234	Both	Mod/High				No	No	No	No	No
540	1234	UC	Mod/High				No	No	No	No	No

Abbreviations: CD = Crohn's Disease: UC = Ulcerative Colitis: Mod/High = Moderate to High; JAK = Janus Kinases

# **Discussion and Conclusions**

- Risk stratification of patients with IBD is not explicitly documented in the EMR.
- Nearly 80% of cases were mod/high risk and 67% of CD cases had perianal disease.
- Ochsner population is nearly 36% Black, whom have been found to have more perianal disease.<sup>9</sup>
- Tools such as IBD PATH can inform PHM of patients diagnosed with IBD, facilitating the identification of potential population level gaps in care for further assessment.
- Results are dependent on the completeness of data uploaded. Incomplete data may result in the underreporting of medications and quality indicators.

1. Yu Y, et al. Prevalence of inflammatory bowel disease in pediatric and adult populations: recent estimates from large national databases in the United States, 2007-2016. Inflamm Bowel

2. Park KT, et al. The cost of inflammatory bowel disease: an initiative from the Crohn's & Colitis Foundation. Inflamm Bowel Dis. 2020;26(1):1-10.

8. Siegel CA, et al. Identifying patients with inflammatory bowel diseases at high vs low risk of complications. Clin Gastroenterol Hepatol. 2020;18(6):1261-1267.