Assessing Disease Severity in Ulcerative Colitis

- Multiple severity scales available
- Mayo Score Combination of patient reported outcomes, PGA, and endoscopy¹

| Variable | Symptoms | Score | Variable | Symptoms | Score | | |
|-----------------|--|-------|-----------|---------------------------------------|-------|--|----------|
| Stool Frequency | 1-2 more bowel movements/day than normal | 0 | Endoscopy | Normal/inactive colitis | 0 | Common Severity Thresholds for Full Score (not validated) ^{1,2} | |
| | | 1 | | Erythema, decreased vascularity | 1 | | |
| | 3-4 more bowel movements/day | 1 | | Friability, marked erythema, erosions | 2 | | |
| | 5-6 more bowel movements/day | 2 | | Ulcerations, severe friability, | 3 | Score | Severity |
| | >6 more bowel movements/day | 3 | | spontaneous bleeding | - | Remission | <2 |
| Rectal Bleeding | None | 0 | PGA | Normal | 0 | Mild | 2-5 |
| | <50% of stools | 1 | | Mild | 1 | Moderate- | 6-12 |
| | >50% of stools | 2 | | Moderate | 2 | to-Severe | |
| | Blood only | 3 | | Severe | 3 | | |

Both Modified and Partial Mayo Scores correlate with full scores²

- Modified Mayo Score no PGA, useful for clinical trials
- Partial Mayo Score no endoscopy, useful in clinical practice

PGA, physician's global assessment.

1. Feurstein JD, et al. *Gastroenterology*. 2020;158(5):1450-61; 2. Naegeli AN, et al. *Crohns Colitis 360*. 2021;3(1):otab007.

Proposed Ulcerative Colitis Activity Index

| Proposed American College of Gastroenterology Ulcerative Colitis Activity Index | | | | | | | | |
|---|---------------|------------------|--------------------|----------------------|--|--|--|--|
| | Remission | Mild | Moderate-to-Severe | Fulminant | | | | |
| Stools (No./d) | Formed stools | <4 | >6 | >10 | | | | |
| Blood in stools | None | Intermittent | Frequent | Continuous | | | | |
| Urgency | None | Mild, occasional | Often | Continuous | | | | |
| Hemoglobin | Normal | Normal | <75% of normal | Transfusion required | | | | |
| ESR | <30 | <30 | >30 | >30 | | | | |
| CRP (mg/L) | Normal | Elevated | Elevated | Elevated | | | | |
| FC (µg/g) | <100-200 | >150-200 | >150-200 | >150-200 | | | | |
| Endoscopy (Mayo score) | 0-1 | 1 | 2-3 | 3 | | | | |
| UCEIS | 0-1 | 2-4 | 5-8 | 7-8 | | | | |

General guides—with the exception of remission, all the factors are not needed for a patient to be considered in a specific category

CRP, C-reactive protein; ESR, erythrocyte sedimentation rate; FC, fecal calprotectin; UCEIS, UC Endoscopic Index of Severity. Rubin DT, et al. *Am J Gastroenterol*. 2019;114(3):384-413.

Assessing Disease Severity in Crohn's Disease

- Crohn's Disease Activity Index (CDAI) is commonly used in clinical trials¹
- Includes multiple parameters, assessed over past 7 days:²
 - Stool pattern
 - Abdominal pain
 - General well-being
 - Complications
 - Finding of abdominal mass
 - Anemia
 - Weight change

| Severity Thresholds ¹ | | | | | |
|----------------------------------|----------|--|--|--|--|
| Score | Severity | | | | |
| Remission | <150 | | | | |
| Mild | 150-220 | | | | |
| Moderate | >220-450 | | | | |
| Severe | >450 | | | | |

Individualized Treatment and Disease Characteristics

Disease Severity

More aggressive therapy recommended for greater severity¹⁻⁴

- Step-up approach not recommended for moderate-to-severe CD or UC
- Biologics or small-molecule therapies recommended for first-line treatment; anti-TNF-α drugs are the most extensively studied

Additional Considerations

1. Feurstein JD, et al. *Gastroenterology*. 2021;160(7):2496-2508; 2. Lichtenstein GR, et al. *Am J Gastroenterol*. 2018;113(4):481-517; 3. Feurstein JD, et al. *Gastroenterology*. 2020;158(5):1450-61; 4. Rubin DT, et al. *Am J Gastroenterol*. 2019;114(3):384-413; 5. Garcia NM, et al. *United European Gastroenterol J*. 2022; 10(10):1121-8.

Previous Treatments

Previous history with biologics influences treatment choice¹

- Vedolizumab associated with greater clinical and/or endoscopic remission in anti-TNF- α naïve patients than in those with previous exposure
- After anti-TNF- α primary or secondary nonresponse, a second anti-TNF- α agent is less likely to achieve clinical remission

AGA recommended first-line therapy for moderate-to-severe IBD

CD – infliximab, adalimumab, or ustekinumab²

UC – infliximab or vedolizumab³

AGA recommended therapy following anti-TNF- α nonresponse for moderate-to-severe IBD

CD – ustekinumab or adalimumab (adalimumab for primary nonresponse only)²

UC – ustekinumab or tofacitinib*³

^{*}Tofacitinib was the only Janus kinase inhibitor available at time of guidelines publication. AGA, American Gastroenterological Association.

^{1.} Bressler B. Therap Adv Gastroenterol. 2023;16:17562848231159452; 2. Feurstein JD, et al. Gastroenterology. 2021;160(7):2496-2508; 3. Feurstein JD, et al. Gastroenterology. 2020;158(5):1450-61.