

## Iowa Department of Human Services

## Request for Prior Authorization SELECT NON-BIOLOGIC AGENTS FOR ULCERATIVE COLITIS



(PLEASE PRINT - ACCURACY IS IMPORTANT)

FAX Completed Form To 1 (877) 733-3195 Provider Help Desk 1 (844) 236-1464

IA Medicaid Member ID #	Patient name	DOB	
Patient address			
Provider NPI	Prescriber name	Phone	
Prescriber address		Fax	
Pharmacy name	Address	Phone	
Prescriber must complete all inform	mation above. It must be legible, correct, and c	omplete or form will be returned.	
Pharmacy NPI	Pharmacy fax	NDC 	
Prior authorization is required for select non-biologicals for ulcerative colitis (UC). Payment for non-preferred select non-biologicals for UC may be considered only for cases in which there is documentation of a previous trial and therapy failure with the preferred agent(s). Payment will be considered under the following conditions:  1) Patient has a diagnosis of moderately to severely active UC; and  2) Request adheres to all FDA approved labeling for indication, including age, dosing, and contraindications; and  3) A documented trial and inadequate response to two preferred conventional therapies (immunomodulators) including aminosalicylates and azathioprine/6-mercaptopurine; and  4) A documented trial and inadequate response with a preferred biological DMARD; and  5) Will not be taken concomitantly with immunomodulators or biologic therapies.  The required trials may be overridden when documented evidence is provided that use of these agents would be medically contraindicated.  Non-Preferred  Zeposia			
Strength Dosa			
	ge Instructions Q	uantity Days Supply	
	ge Instructions Q	uantity Days Supply	
Diagnosis:			
Diagnosis:  Will medication be used in c			
Diagnosis:	ombination with immunomodulators o		
Diagnosis:  Will medication be used in c  Yes No  Trial Documentation:  Preferred Conventional There	ombination with immunomodulators o	r biologic therapies?	
Diagnosis:  Will medication be used in c  ☐ Yes ☐ No  Trial Documentation:  Preferred Conventional There  Trial 1: Name/Dose:	ombination with immunomodulators o	r biologic therapies? Trial Dates:	
Diagnosis:  Will medication be used in c  ☐ Yes ☐ No  Trial Documentation:  Preferred Conventional There  Trial 1: Name/Dose:	ombination with immunomodulators o	r biologic therapies? Trial Dates:	
Diagnosis:  Will medication be used in c  ☐ Yes ☐ No  Trial Documentation:  Preferred Conventional Ther  Trial 1: Name/Dose:  Failure reason:	ombination with immunomodulators o	r biologic therapies? Trial Dates:	
Diagnosis:  Will medication be used in c  ☐ Yes ☐ No  Trial Documentation:  Preferred Conventional Ther  Trial 1: Name/Dose:  Failure reason:	ombination with immunomodulators o	r biologic therapies? Trial Dates:	

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Preferred Biological DMARD:			
Name/Dose:T	rial Dates:		
Failure reason:			
Medical or contraindication reason to override trial requirements:			
Attach lab results and other documentation as necessary.			
Prescriber signature (Must match prescriber listed above.)	Date of submission		

**IMPORTANT NOTE:** In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Human Services, that the member continues to be eligible for Medicaid.

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