SENTARA COMMUNITY PLAN (MEDICAID)

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions:</u> The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; <u>fax to 1-844-305-2331</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If information provided is not</u> complete, correct, or legible, authorization can be delayed.

Drug Requested: Entyvio® IV (vedolizumab) (J3380) (Medical)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.		
Member Name:		
Member Sentara #:	Date of Birth:	
Prescriber Name:		
Prescriber Signature:	Date:	
Office Contact Name:		
Phone Number:	Fax Number:	
NPI #:		
DRUG INFORMATION: Authorization may be		
Drug Name/Form/Strength:		
Dosing Schedule:	Length of Therapy:	
Diagnosis:	ICD Code, if applicable:	
Weight (if applicable):	Date weight obtained:	
Standard Review. In checking this box, the timeframe does not jeopardize the life or health of the memb or the member's ability to regain maximum function and would not subject the member to severe pain.		
• Entyvio 300 mg/mL solution; 1 vial = 300 billabl	e units	
For Crohn's disease or Ulcerative Colitis: IV – weeks) and then 300mg (1 vial) every 8 weeks thereafter who show no evidence of therapeutic benefit by week 14	the induction period. Discontinue therapy in patients	
Off-label dosing:		

(Continued on next page)

Please submit literature and progress notes for off-label dosing.

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

DIAGNOSIS: Check diagnosis that applies.			
	Crohn's Disease		Ulcerative Colitis:
Ţ			men of oral corticosteroids (moderate to severe CD or costeroids (severe and fulminant CD or UC or failure to
(☐ Member has a trial and failure of a compliant regimen of azathioprine or mercaptopurine for at least three (3) consecutive months		
(Member has a trial and failure of a compliant consecutive months	regi	gimen of parenteral methotrexate for at least three (3)
(☐ Member has tried and failed: ☐ Humi	ra®	AND Infliximab
Medication being provided by: Please check applicable box below.			
	Location/site of drug administration:		
NPI or DEA # of administering location:			
	<u>OR</u>		
	Specialty Pharmacy – PropriumRx		

For urgent reviews: Practitioner should call Sentara Health Pre-Authorization Department if they believe a standard review would subject the member to adverse health consequences. Sentara Health's definition of urgent is a lack of treatment that could seriously jeopardize the life or health of the member or the member's ability to regain maximum function.

**Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *