AvMed

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: 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 1-305-671-0200</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If the information provided is not complete, correct, or legible, the authorization process can be delayed.

Drug Requested: Actemra[®] (tocilizumab) SubQ Giant Cell Arteritis (GCA) (self-administered)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.					
Member Name:					
Member AvMed #:					
Prescriber Name:					
Prescriber Signature:					
Office Contact Name:					
	Fax Number:				
DEA OR NPI #:					
DRUG INFORMATION: Authorization may be delayed if incomplete.					
Drug Form/Strength:					
Dosing Schedule:	Length of Therapy:				
Diagnosis:	ICD Code, if applicable:				
Weight:	Date:				
<u>Recommended Dose for Actemra[®] for adult members with GCA</u> - 162 mg given once every week as a subcutaneous injection, in combination with a tapering course of glucocorticoids					
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 dated within 60 days, must be provided or request may be denied.					

• Must be prescribed by or in consultation with (check box below that applies):

Neurologist	Rheumatologist	Ophthalmologist
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□ Member has diagnosis of Giant Cell Arteritis (GCA)

AND

□ Member is at least 50 years of age

AND

□ Member has ESR >30mm/hour <u>OR</u> CRP> 1 mg/dL currently on prednisone

AND

- □ Member had trial and failure of <u>ONE</u> of the following:
 - □ 40mg Prednisolone daily for 4 weeks
 - □ 80mg Prednisolone daily if eye symptoms for 4 weeks

OR

- Member has a contraindication to prednisolone and documentation that GI BLEED occurred within the last 30 days has been submitted (medical chart notes must be attached) <u>AND</u> member has <u>one</u> of the following (labs must be submitted):
 - □ ESR >50mm/hour not currently on prednisolone

OR

□ CRP> 2.49 mg/dL not currently on prednisolone

AND

Medical chart notes documenting the following <u>MUST</u> be submitted:

- □ Unequivocal cranial symptoms of GCA new-onset at least <u>TWO</u> of the following features must be present:
 - □ Localized headache, scalp tenderness, temporal artery tenderness, decrease pulsation, ischemiarelated vision loss, or otherwise unexplained mouth or jaw pain upon mastication

AND

At least **ONE** of the following **MUST** be submitted for documentation:

□ Temporal artery biopsy revealing features of GCA must be submitted documenting at least <u>TWO (2)</u> of the following:

Granulomatous inflammation of the blood vessel wall	Disruption and fragmentation of internal elastic lamina	Giant cells
Proliferation of the intima with associated occlusion of the lumen	The healed stage reveals collagenous thickening of the vessel wall and the artery is transformed into a fibrous cord	

OR

(Continued on next page)

- Magnetic resonance angiography (MRA), Computed tomography angiography (CTA), or Positron emission tomography-computed tomography angiography (PET-CTA) must_be submitted to document the following:
 - □ Evidence of large-vessel vasculitis by angiography or cross-sectional imaging study

Medication being provided by (check box below that applies):

OR

Physician's office

Specialty Pharmacy - PropriumRx

Not all drugs may be covered under every Plan

If a drug is non-formulary on a Plan, documentation of medical necessity will be required. **Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.** *<u>Previous therapies will be verified through pharmacy paid claims or submitted chart notes.</u>*