

# SENTARA HEALTH PLANS

## PHARMACY/MEDICAL 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; **fax to 1-800-750-9692.** No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If information provided is not complete, correct, or legible, authorization can be delayed.**

**Drug Requested:** Actemra<sup>®</sup> (tocilizumab) - Giant Cell Arteritis (GCA) (self-administered)  
SubQ

**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: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

**DRUG INFORMATION:** Authorization may be delayed if incomplete.

Drug Form/Strength/Quantity: \_\_\_\_\_

Dosing Schedule: \_\_\_\_\_ Length of Therapy: \_\_\_\_\_

Diagnosis: \_\_\_\_\_ ICD Code, if applicable: \_\_\_\_\_

Weight: \_\_\_\_\_ Date: \_\_\_\_\_

**Recommended Dose for Actemra<sup>®</sup> for adult members with GCA** - 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

Member has diagnosis of Giant Cell Arteritis (GCA)

**AND**

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- Member is at least 50 years of age
- AND**
- Member has ESR >30mm/hour **OR** CRP> 1 mg/dL currently on prednisone

**AND**

- Member had trial and failure of **ONE** 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**) **AND** member has **one** 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 MUST BE SUBMITTED:**

- Unequivocal cranial symptoms of GCA new-onset - at least **TWO** of the following features must be present:
  - Localized headache, scalp tenderness, temporal artery tenderness, decrease pulsation, ischemia-related 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 **TWO (2)** of the following:

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

**OR**

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- 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):**

- Physician's office                      **OR**                       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.\*\**

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