

SENTARA COMMUNITY PLAN (MEDICAID)

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; **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 the information provided is not complete, correct, or legible, the authorization process can be delayed.**

Drug Requested: Orencia® SQ (abatacept) (Pharmacy)

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 delayed if incomplete.

Drug Name/Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight (if applicable): _____ Date weight obtained: _____

DIAGNOSIS	Recommended Dose
<ul style="list-style-type: none">Moderate to severe Active Rheumatoid Arthritis (RA)Psoriatic Arthritis (PsA)	SUBCUTANEOUS <ul style="list-style-type: none">125 mg once a week (4 syringe/28 days) <u>Pediatric dosing:</u> <ul style="list-style-type: none">Weighing 10 kg to less than 25 kg: 50 mg once a week (1 syringe/28 days)Weighing 25 kg to less than 50 kg: 87.5 mg once a week (1 syringe/28 days)Weighing 50 kg or greater: 125 mg once a week (4 syringes/28 days)

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DIAGNOSIS	Recommended Dose
Juvenile Idiopathic Arthritis (pJIA) in members 2 years and older	SUBCUTANEOUS <ul style="list-style-type: none"> • Weighing greater 50 kg: 125 mg once a week (4 syringes/28 days) • Weighing 25 kg to less than 50 kg: 87.5 mg (Four syringes of 87.5mg/28days) once a week • Weighing 10kg to 25 kg: 50 mg once a week (Four syringes of 50 mg/28 days)

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 applicable diagnosis below

Moderate to severe Active Rheumatoid Arthritis (RA)

- Member is 18 years of age or older
- Member has been diagnosed with moderate to severe rheumatoid arthritis
- Trial and failure of, contraindication, or adverse reaction to methotrexate
- Trial and failure of at least **ONE (1)** other **DMARD therapy** including, but not limited to: (**check each tried**)

<input type="checkbox"/> auranofin	<input type="checkbox"/> sulfasalazine
<input type="checkbox"/> azathioprine	<input type="checkbox"/> leflunomide
<input type="checkbox"/> hydroxychloroquine	<input type="checkbox"/> other: _____

- Trial and failure of **BOTH** of the preferred drugs below:

<input type="checkbox"/> adalimumab-adbm (Boehringer Ingelheim) OR Hadlima [®] (adalimumab-bwwd)	<input type="checkbox"/> Enbrel [®]
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Polyarticular Juvenile Idiopathic Arthritis (pJIA)

- Member is 2 years of age or older
- Member has been diagnosed with moderate to severe active Polyarticular Juvenile Idiopathic Arthritis (pJIA)
- Trial and failure of **BOTH** of the preferred drugs below:

<input type="checkbox"/> adalimumab-adbm (Boehringer Ingelheim) OR Hadlima [®] (adalimumab-bwwd)	<input type="checkbox"/> Enbrel [®]
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