

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: Stelara™ SQ (ustekinumab) (PHARMACY BENEFIT ONLY)
(Stelara SQ therapy is **Self-Administered** by member)

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: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight: _____ Date: _____

Recommended Dose:

Indication	Dosage:
Adults with Moderate to Severe Chronic Plaque Psoriasis	Weight <ul style="list-style-type: none">• Less than or = 100 kg Initial (two 45 mg prefilled syringe/ 28 days) then continue with one 45 mg prefilled syringe/84 days• Great than 100 kg Two 90 mg administered prefilled syringe/ 28 days then, one 90 mg administered prefilled syringe/ 84 days

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Indication	Dosage:
Adolescent patients (6 years or older) with Moderate to Severe Chronic Plaque Psoriasis	<u>6 years and older SQ</u> <ul style="list-style-type: none"> • Less than or = 60 kg (0.75 mg/kg) dosing down to 15 kg • ≥ 60 to ≤ 100 kg • Initial (two 45 mg prefilled syringe/ 28 days) then continue with one 45 mg prefilled syringe/ 84 days • Greater than 100 kg Two 90 mg administered prefilled syringe/ 28 days, then one 90 mg
Active Psoriatic Arthritis	<ul style="list-style-type: none"> • Two 45 mg prefilled syringe/28 days; then continue with one 45 mg prefilled syringe/ 84 days
Moderately to Severe Active Crohn's Disease	<ul style="list-style-type: none"> • A single intravenous infusion using weight-based dosing • Up to 55 kg 260 mg (2 vials) • Greater than 55 kg to 85 kg 390 mg (3 vials) • Greater than 85 kg 520 mg (4 vials)
Adult patient with Moderately to Severely Active Ulcerative Colitis	<ul style="list-style-type: none"> • Intravenous dose; then every 8 weeks thereafter (1 vial per 56 days) • A subcutaneous 90 mg dose 8 weeks after the initial (90 mg prefilled syringe/56 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

Prescriber is a: Dermatologist Rheumatologist Gastroenterologist

Diagnosis: Check below diagnosis that applies to qualify for approval or authorization may be delayed.

Active Psoriatic Arthritis

Member tried and failed at least **one DMARD** (Check each tried):

<input type="checkbox"/> methotrexate	<input type="checkbox"/> sulfasalazine	<input type="checkbox"/> azathioprine
<input type="checkbox"/> leflunomide	<input type="checkbox"/> auranofin	<input type="checkbox"/> hydroxychloroquine

AND

Patient has tried and failed **TWO (2)** of the following biologics:

<input type="checkbox"/> Humira®	<input type="checkbox"/> Enbrel®	<input type="checkbox"/> Infliximab
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Moderate to Severe Chronic Plaque Psoriasis

- Member is 6 years or older with moderate to severe plaque psoriasis who are candidates for phototherapy or systemic therapy.

AND

- Trial and failure of, contraindication, or adverse reaction to methotrexate

AND

- Trial and failure of **TWO (2)** of the **PREFERRED** drugs below:

<input type="checkbox"/> Humira®	<input type="checkbox"/> Enbrel®	<input type="checkbox"/> Infliximab
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DIAGNOSIS: Check diagnosis that applies.

<input type="checkbox"/> Crohn's Disease	<input type="checkbox"/> Ulcerative Colitis:
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- Member has trial and failure of a compliant regimen of oral corticosteroids (budesonide 9mg daily for 8 weeks) or high dose steroids (40-60 mg prednisone) (moderate to severe CD) unless contraindicated or intravenous corticosteroids (severe and fulminant CD or failure to respond to oral corticosteroids)

AND

- Member tried and failed **at least ONE previous 5-Aminosalicylates or immunomodulators therapy below:**

<input type="checkbox"/> methotrexate	<input type="checkbox"/> azathioprine	<input type="checkbox"/> auranofin
<input type="checkbox"/> sulfasalazine	<input type="checkbox"/> oral aminosalicylates	<input type="checkbox"/> leflunomide
<input type="checkbox"/> 6-mercaptopurine	<input type="checkbox"/> Apriso®	<input type="checkbox"/> balsalazide
<input type="checkbox"/> Pentasa®		

AND

- Member has tried and failed both:

<input type="checkbox"/> Humira®	<input type="checkbox"/> Infliximab
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Medication being provided by a Specialty Pharmacy - PropriumRx

****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.****
Previous therap-ies will be verified through pharmacy paid claims or submitted chart notes.