

# 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:** Cibinco<sup>TM</sup> (abrocitinib) **(Non-Preferred)**

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

### **Recommended Dosage:**

- 100 mg orally once daily
- 200 mg orally once daily if not responding to 100 mg daily

**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: Refractory, Moderate-to-Severe Atopic Dermatitis**

**Length of authorization: 12 months.**

☐ Member is  $\geq 12$  years of age

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- ❑ Diagnosis of refractory, moderate-to-severe atopic dermatitis
- ❑ Prior documented trial and failure (or contraindication) **BOTH** of the following:
  - ❑ One (1) topical corticosteroid of medium to high potency (e.g., mometasone, fluocinolone)
  - ❑ One (1) topical calcineurin inhibitors (tacrolimus or pimecrolimus)
- ❑ Inadequate response to a 3-month minimum trial of at least one immunosuppressive systemic agent (e.g., cyclosporine, azathioprine, methotrexate, mycophenolate mofetil, etc.)
- ❑ Inadequate response (or is not a candidate) to a 3-month minimum trial of phototherapy (e.g., psoralens with UVA light [PUVA], UVB, etc.) provided member has reasonable access to photo treatment
- ❑ Prescriber attestation that Cibinqo will not be used in combination with other JAK inhibitors, biologic immunomodulators, or with other immunosuppressants
- ❑ Trial and failure of Dupixent<sup>®</sup> (dupilumab)

<b>Medication being provided by Specialty Pharmacy - PropriumRx</b>
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***\*\*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.\****