

# 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: (Check applicable drug below)

**Lampit<sup>®</sup>** (nifurtimox) tablets

**benznidazole** tablets

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

### Quantity Limits based on age and weight:

- **Lampit<sup>®</sup>:** Maximum of 10 to 20 mg/kg/day for those weighing 2.5 to < 40kg, and 8 to 10 mg/kg/day for those weighing ≥ 40kg, in 3 divided doses for 60 days
- **benznidazole:** Maximum of 5 to 8 mg/kg/day in 2 divided doses, administered every 12 hours for 60 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.

### Approval Length: 60 Days

- For Lampit<sup>®</sup>: Member is < 18 years of age
- For benznidazole: Member is 2-12 years of age

(Continued on next page)

**AND**

- ❑ Medication is prescribed by an infectious disease specialist

**AND**

- ❑ Confirmation of Chagas disease was made through positive identification by microscopy or serological assay of *Trypanosoma crusi* (**\*coverage excluded for other species of *Trypanosoma*\***)(**lab results must be submitted**)

**AND**

- ❑ For females of reproductive potential: Pregnancy has been evaluated prior to treatment, will be monitored during treatment, and contraception is made available due to potential for teratogenicity of these agents

**AND**

- ❑ Provider attests that monitoring of blood cell counts will be done at baseline and during therapy with nifurtimox (Lampit<sup>®</sup>) or benznidazole

**AND**

- ❑ For Lampit<sup>®</sup>: Provider attests hepatic and renal monitoring will be done at baseline and during therapy

**Medication being provided by a Specialty Pharmacy - PropriumRx**

***\*\*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.\****