

# AvMed

## 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 1-305-671-0200.** 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: Repository Corticotropin Medications - INFANTILE SPASMS (IS)

<u>PREFERRED</u>	<u>NON-PREFERRED</u>
<input type="checkbox"/> <b>Purified Cortrophin™ Gel</b> (repository corticotropin)	<input type="checkbox"/> <b>HP Acthar® Gel</b> (repository corticotropin) *Member must have tried and failed preferred Purified Cortrophin™ Gel and meet all applicable PA criteria below

**\*\*Acthar Gel single-dose pre-filled SelfJect injector is for subcutaneous administration by adults only.\*\***

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

**Note:** (Neurology 2012;78:1974-1976) Class I study showed similar efficacy between low-dose (20-30 IU) and high dose (150 IU/m<sup>2</sup>) natural ACTH. Low dose ACTH should be considered as an alternative to high dose ACTH for treatment of infantile spasms. (Level B).

**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.

(Continued on next page)

- Prescriber **MUST** be a Neurologist
- Member **MUST** have a documented diagnosis of Infantile Spasms
- Approval will only be granted for a **MAXIMUM** of **30 days only** due to similar adverse effects of corticosteroids. After 2 weeks of treatment, dosing should be gradually tapered and discontinued over a 2-week period. The following is one **suggested** tapering schedule:
  - 30 U/m<sup>2</sup> in the morning for 3 days; 15 U/m<sup>2</sup> in the morning for 3 days; 10 U/m<sup>2</sup> in the morning for 3 days; and 10 U/m<sup>2</sup> every other morning for 6 days.
- Complete the regimen below (**repository corticotropin is supplied as 5 mL multi-dose vial containing 80 USP Units per mL**):

Approval will be a **MAXIMUM** of **30 days only** (combined inpatient and outpatient time period)

<u>Initial Dose Schedule</u>	<u>Approval at Outpatient pharmacy will be based on volume needed at discharge from hospital</u>	
75 U/m <sup>2</sup> BID x _____ days	TOTAL _____ mL x _____ # days (max 29 days)	
<u>Taper Dose Schedule</u>		<u>BODY SURFACE AREA BSA</u>
30 U/m <sup>2</sup> QD x _____ days	_____ mL x _____ days	Weight: _____ kg
15 U/m <sup>2</sup> QD x _____ days	_____ mL x _____ days	Height/Length: _____ in.
10 U/m <sup>2</sup> QD x _____ days	_____ mL x _____ days	Calculated BSA: _____ m <sup>2</sup>
10 U/m <sup>2</sup> QOD x _____ days	_____ mL x _____ days	

TOTAL Number of vials needed: \_\_\_\_\_ /days (max 29 days)

**Medication being provided by Specialty Pharmacy – Proprium Rx**

*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.\**