

# OPTIMA HEALTH PLAN

## PHARMACY/MEDICAL PRIOR AUTHORIZATION 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. Incomplete form will delay authorization process.

**Drug Requested:**                    **Acthar® HP (Corticotropin) - INFANTILE SPASMS (IS)**

**DRUG INFORMATION:** Complete all information below or authorization process will be delayed.

**Drug Form/Strength/Month:** \_\_\_\_\_

**Dosing Schedule:** \_\_\_\_\_                    **Length of Therapy:** \_\_\_\_\_

**Diagnosis:** \_\_\_\_\_                    **ICD Code, if applicable:** \_\_\_\_\_

**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:** The criteria below **must** be met to qualify or authorization process will be delayed.

- Prescriber **MUST** be a Neurologist

**AND**

- Patient **MUST** have a documented diagnosis of Infantile Spasms

**AND**

- Approval will only be granted for a **MAXIMUM** of **30 days only** due to similar adverse effect 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 (*HP Acthar gel is supplied as 5mL multidose 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> <b>BID</b> 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> <b>QD</b> x _____ days  | _____ mL x _____ days  | <b>WEIGHT:</b> _____ kg              |
| 15 U/m <sup>2</sup> <b>QD</b> x _____ days  | _____ mL x _____ days  | Height/Length: _____ in.             |
| 10 U/m <sup>2</sup> <b>QD</b> x _____ days  | _____ mL x _____ days  | Calculated BSA: _____ m <sup>2</sup> |
| 10 U/m <sup>2</sup> <b>QOD</b> x _____ days | _____ mL x _____ days  |                                      |

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

(signature on next page)

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.\***

Patient Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

**DEA OR NPI #:** \_\_\_\_\_

**\*Approved by Pharmacy and Therapeutics Committee: 2/21/2008**

**UPDATED:** 6/2/2011; 8/11/2011; 10/1/2012; 8/19/2014; 10/31/2014; 4/3/2015; 5/23/2015; 6/8/2015; 12/22/2015; 6/15/2016; 8/25/2016; 9/22/2016; 12/11/2016; 5/25/2017; 7/30/2017; **6/22/2018**