

Pharmacy Prior Authorization Form Fax completed form to: 877.974.4411 toll free, or 616.942.8206 This form applies to: Medicaid ☐ **Urgent** (life threatening) ☐ **Non-Urgent** (standard review) This request is: Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function.. H.P. Acthar[®] (corticotropin) Member First Name: _____ Gender: _____ Last Name: #:____Primary Care Physician:_____ Requesting Provider: Provider Address: Provider NPI: Contact Name: Provider Signature: **Product and Billing Information** □ New Request □ Continuation Request Start date (or date of next dose): Drug product: ☐ H.P. Acthar Gel 80 units/mL inj. Date of last dose (if applicable):

Additional Information

H.P. Acthar Gel is covered for infantile spasms (West syndrome) when precertification requirements are met. H.P. Acthar Gel is not considered medically necessary for corticosteroid-responsive conditions because it has not been proven to be more effective than corticosteroids for these conditions.

Dose and frequency:

H.P. Acthar Gel is not considered medically necessary for all other indications including, but not limited to:

- 1. Acute exacerbations of multiple sclerosis
- 2. Rheumatic disorders (psoriatic arthritis, rheumatoid arthritis, ankylosing spondylitis)
- 3. Collagen diseases (systemic lupus erythematosus, systemic dermatomyositis)
- 4. Dermatologic diseases (severe erythema multiforme, Stevens-Johnson syndrome)
- 5. Allergic states (serum sickness)
- 6. Ophthalmic diseases (keratitis, iritis, iridocyclitis, uveitis, choroiditis, optic neuritis, chorioretinitis, anterior segment inflammation)
- 7. Respiratory diseases (symptomatic sarcoidosis)
- 8. Edematous state

Precertification Requirements

Before this drug is covered, the patient must meet the requirements for one of the following conditions:

1. For a diagnosis of infantile spasms for patients under 2 years of age, H.P. Acthar Gel is authorized up to a dose of 75 units/m² twice daily for two weeks, followed by a tapering schedule for an additional two weeks

Note: Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMS-accepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, the dosing of the drug, or the route of administration to be used for the identified indication.



	New request Priority Health Precertification Documentation	
A.	What is the patient's diagnosis?	
	☐ Infantile spasms ☐ Other – rationale for use:	