

## PHARMACY POLICY STATEMENT North Carolina Marketplace

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| <b>DRUG NAME</b>        | <b>Acthar Gel (repository corticotropin injection)</b> |
| BILLING CODE            | J0800  |
| BENEFIT TYPE            | Medical or Pharmacy                                    |
| SITE OF SERVICE ALLOWED | Home/Office  |
| STATUS                  | Prior Authorization Required                           |

Acthar is a corticotropin initially approved by the FDA in 1952. It is indicated as monotherapy for the treatment of infantile spasms in infants and children under 2 years of age. The mechanism of action of Acthar Gel in the treatment of infantile spasms is unknown. Acthar Gel and endogenous ACTH stimulate the adrenal cortex to secrete cortisol, corticosterone, aldosterone, and a number of weakly androgenic substances. Prolonged administration of large doses of Acthar Gel induces hyperplasia and hypertrophy of the adrenal cortex and continuous high output of cortisol, corticosterone and weak androgens. The release of endogenous ACTH is under the influence of the nervous system via the regulatory hormone released from the hypothalamus and by a negative corticosteroid feedback mechanism. Elevated plasma cortisol suppresses ACTH release. Acthar Gel is also reported to bind to melanocortin receptors. The trophic effects of endogenous ACTH and Acthar Gel on the adrenal cortex are not well understood beyond the fact that they appear to be mediated by cyclic AMP.

Acthar Gel (repository corticotropin injection) will be considered for coverage when the following criteria are met:

### Infantile Spasms (West syndrome, X-linked infantile spasms syndrome)

For **initial** authorization:

1. Member is an infant or a child under 2 years of age; AND
2. Medication must be prescribed by a pediatric neurologist or an epileptologist; AND
3. Member has documented diagnosis of infantile spasms in chart notes; AND
4. Member's body surface area (BSA, m<sup>2</sup>) or height and weight have been provided to determine the appropriate dosage.
5. Medication is used as monotherapy; AND
6. **Dosage allowed:** The recommended regimen is a maximum daily dose of 150 U/m<sup>2</sup> (divided into twice daily injections of 75 U/m<sup>2</sup>) for 2 weeks. After 2 weeks of treatment, dosing should be gradually tapered and discontinued over a 2-week period. The dosing calculator is available on Acthar's website. Quantity Limit: 3 vials (15 mL total) per 28 days

***If all the above requirements are met, the medication will be approved for 1 month.***



For **reauthorization**:

1. Member must be under 2 years of age; AND
2. Chart notes demonstrate clinical benefit from the initial use of medication (e.g., suppression of spasm symptoms); AND
3. Member experienced a relapse in spasm symptoms after Acthar was discontinued.

***If all the above requirements are met, the medication will be approved for an additional month.***

**CareSource considers Acthar Gel (repository corticotropin injection) not medically necessary for the treatment of conditions that are not listed in this document. For any other indication, please refer to the Off-Label policy.**

| DATE       | ACTION/DESCRIPTION   |
|------------|--|
| 10/08/2018 | New policy for H.P.Acthar created. Policy placed in the new format.  |
| 01/22/2021 | Changed name to Acthar. Increased the quantity limit to 3 vials (15 mL) per 28 days. Adjusted specialist name. Added that BSA or height/weight must be provided to calculate quantity. Reworded reauth requirement to be more specific. Added member must be under 2 years of age for reauth. Added that member must experience relapse in spasm symptoms after Acthar was discontinued. Updated references. |
| 05/05/2022 | Transferred to new format. Updated references. Added medication must be used as monotherapy.   |

References:

1. H.P. Acthar Gel [package insert]. Hazelwood, MO: Mallinckrodt ARD Inc.; October 2021.
2. AAN/CNS evidence-based guideline update on medical treatment of infantile spasms. *Neurology* 2012; 78 (24): 1974 – 80. doi: 10.1212/WNL.0b013e318259e2cf.
3. Wilmshurst JM, Gaillard WD, Vinayan KP, et al. Summary of recommendations for the management of infantile seizures: Task Force Report for the ILAE Commission of Pediatrics. *Epilepsia*. 2015;56(8):1185-1197. doi:10.1111/epi.13057.
4. Nelson GR. Management of infantile spasms. *Transl Pediatr*. 2015;4(4):260-270. doi:10.3978/j.issn.2224-4336.2015.09.01.
5. Gold Standard, Inc. Corticotropin ACTH. Clinical Pharmacology [database online]. Tampa, FL: Gold Standard, Inc; 2012. Available from: <http://www.clinicalpharmacology.com>.
6. Management and prognosis of infantile spasms. Daniel G Glaze. UpToDate [online database]. Available from: <http://www.uptodate.com>
7. Milanese C, La Mantia L, Salmaggi A, et al. Double-blind randomized trial of ACTH versus dexamethasone versus methylprednisolone in multiple sclerosis bouts. Clinical, cerebrospinal fluid and neurophysiological results. *Eur Neurol*. 1989; 29 (1): 10 – 14.
8. Thompson AJ, Kennard C, Swash M, et al. Relative efficacy of intravenous methylprednisolone and ACTH in the treatment of acute relapse in MS. *Neurology*. 1989; 39 (7): 969 – 971.

Effective date: 01/01/2023

Revised date: 05/05/2022