



DRUG	H.P. Acthar® Gel (repository corticotrophin injection)
Policy#	22100
INDICATIONS	<ol> <li>H.P. Acthar Gel is indicated for diagnostic testing of adrenocortical function</li> <li>H.P. Acthar Gel has limited therapeutic value in those conditions responsive to corticosteroid therapy; in such cases, corticosteroid therapy is considered to be the treatment of choice. H.P. Acthar Gel may be employed in the following disorders:         <ul> <li>a. Nervous system diseases: Acute exacerbations of multiple sclerosis</li> <li>b. Rheumatic disorders: As adjunctive therapy for short-term administration (to tide patient over an acute episode or exacerbation) in:</li></ul></li></ol>
CRITERIA	<ul> <li>ConnectiCare will consider H.P. Acthar to be medically necessary in members</li> <li>With chart note documentation of any of the above conditions (listed in 2a-2h above) that have intolerance or treatment failure of at least two other standard pharmacologic therapy including corticosteroid therapy, AND</li> <li>Prescribing provider is a specialist in the disease state being managed</li> </ul>
	Some off-label use may be medically appropriate and rational in certain circumstances. Off-label drug use will be reviewed for evidence of therapeutic value according to the following criteria:





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	<ol> <li>The drug is FDA-approved</li> <li>The member has tried and failed established FDA approved and/or clinical guideline recommended therapy unless contraindicated</li> <li>Phase III* FDA clinical studies to support the non-FDA approved use.</li> <li>A. The drug is recognized for treatment of the requested indication in one of the standard reference compendia         <ul> <li>American Hospital Formulary Service – Drug Information (AHFS-DI)</li> <li>Thomson Micromedex DrugDex</li> <li>Clinical Pharmacology (Gold Standard)</li> <li>National Comprehensive Cancer Network (NCCN)</li> <li>Facts &amp; Comparisons</li> </ul> </li> </ol>
	B. In the absence of being listed in above named sources, a minimum of at least two articles from major peer-reviewed journals (from the United States or great Britain) which supports the proposed use for the specific medical condition as safe and effective. Note: ConnectiCare requires prescribers to submit clinical documentation supporting the drug's effectiveness in treating the intended indication.
	*In Phase III trials, the experimental study drug or treatment is given to large groups of people (1,000-3,000) to confirm its effectiveness, monitor side effects, compare it to commonly used treatments and collect information that will allow the experimental drug or treatment to be used safely.
LIMITATIONS	For members approved for therapy, there is a maximum of 1 vial dispensed per prescription
REFERENCES	1. H. P. Athcar Gel full prescribing information. Union City, CA. Questcor Pharmaceuticals.
P&T REVIEW HISTORY	9/07,6/08, 9/09, 9/10, 12/11, 10/12, 10/13, 10/14, 11/15, 5/16, 5/17, 5/18
REVISION RECORD	11/15, 3/16, 5/17, 5/18