



H.P. Acthar[®] GEL
(repository corticotropin injection) 80 U/mL

SENT PRESCRIPTION DIRECTLY TO SPECIALTY PHARMACY.
PLEASE ENROLL PATIENT IN HUB SERVICES.

PHARMACY NAME: _____

FAX: 1-877-937-2284

Acthar Start Form

Please complete Start Form and fax toll-free
TEL: 1-888-435-2284
Monday through Friday (8:00 AM to 9:00 PM ET)
Saturday (9:00 AM to 2:00 PM ET)

1. PATIENT INFORMATION Patient has been notified of referral YES NO

PATIENT FIRST NAME	PATIENT MIDDLE INITIAL	PATIENT LAST NAME	DATE OF BIRTH	GENDER
HOME ADDRESS		CITY	STATE	ZIP
SHIPPING ADDRESS (IF NOT HOME ADDRESS)	CARE OF (IF NOT ADDRESSED TO PATIENT)	CITY	STATE	ZIP
HOME PHONE	MOBILE	<input type="checkbox"/> OK TO TEXT	BEST TIME TO CALL	
EMAIL ADDRESS	PREFERRED LANGUAGE IF NOT ENGLISH			
ALTERNATIVE CONTACT NAME	RELATIONSHIP TO PATIENT	TELEPHONE		

2. INSURANCE INFORMATION (PLEASE INCLUDE COPIES OF CARDS)

PHARMACY BENEFITS	SUBSCRIBER ID #	GROUP #	TEL #
PRIMARY MEDICAL INSURANCE	SUBSCRIBER ID #	GROUP #	TEL #

3. HEALTHCARE PROVIDER (HCP) INFORMATION

HCP FIRST NAME	HCP LAST NAME	HCP MIDDLE INITIAL	NPI #	GROUP NPI # (IF APPLICABLE)	STATE LICENSE #
SPECIALTY: <input type="checkbox"/> PULMONOLOGY <input type="checkbox"/> RHEUMATOLOGY <input type="checkbox"/> OPHTHALMOLOGY <input type="checkbox"/> OTHER (PLEASE INDICATE) _____					
FACILITY NAME	TELEPHONE	FAX			
ADDRESS	CITY	STATE	ZIP		
OFFICE CONTACT NAME	CONTACT TELEPHONE	MOBILE NUMBER	EMAIL ADDRESS		
PREFERRED METHOD OF COMMUNICATION: <input type="checkbox"/> OFFICE PHONE <input type="checkbox"/> MOBILE PHONE <input type="checkbox"/> FAX <input type="checkbox"/> EMAIL <input type="checkbox"/> TEXT <input type="checkbox"/> NO PREFERENCE					

4. PRESCRIPTION: H.P. ACTHAR[®] GEL NDC# 63004-8710-1 5 mL multidose vial containing 80 USP units per mL

PRIMARY DIAGNOSIS CODES ON PAGE 2, SECTION 6

ICD-10 CODE: _____

DOSE: _____ UNITS mL | ROUTE OF ADMINISTRATION: INTRAMUSCULAR SUBCUTANEOUS | SCHEDULE/FREQUENCY: _____ | QUANTITY OF 5 mL MULTIDOSE VIALS: _____ REFILLS: _____

ALLERGIES _____ ADDITIONAL SPECIAL INSTRUCTIONS, TITRATION OR TAPER DOSE, IF APPLICABLE: _____

IDENTIFYING SUPPLIES IS MANDATORY FOR A COMPLETE PRESCRIPTION:

SYRINGE SIZE: 1 mL 3 mL Other size _____ QUANTITY: 1 box of 100 Other: _____

NEEDLE SIZE FOR DRAWING: 18 G needle, 1" Other: _____ QUANTITY: 1 box of 100 Other: _____

NEEDLE SIZE FOR INJECTION: 25 G needle, 1" (Intramuscular) 25 G needle, 5/8" (Subcutaneous) Other: _____ QUANTITY: 1 box of 100 Other: _____

SUPPLY REFILLS: _____ SHARPS CONTAINER: _____

ACTHAR INJECTION TRAINING SERVICES

By initialing here (original required), I request that company-funded Acthar Injection Training Services be arranged for my patient. I understand that Acthar Injection Training Services are for one instruction visit only and NOT a home health nursing service. I also understand that all reasonable efforts will be made to schedule the Acthar Injection Training Services visit within 24 hours of the patient's receipt of drug shipment. Patients can contact their Case Manager at any time about Injection Training.

HEALTHCARE PROVIDER'S INITIALS _____ DATE _____

5. PRESCRIPTION, CONSENT, AND STATEMENT OF MEDICAL NECESSITY: HCP SIGNATURE REQUIRED

I certify that H.P. Acthar[®] Gel is medically necessary for this patient and that I have reviewed this therapy with the patient and will be monitoring the patient's treatment. I verify that the patient and healthcare provider information on this enrollment form was completed by me or at my direction and that the information contained herein is complete and accurate to the best of my knowledge. I understand that I must comply with my practicing state's specific prescription requirements, such as e-prescribing, state-specific prescription form, fax language, etc. Noncompliance with state-specific requirements could result in outreach to me by the dispensing pharmacy.

I authorize United BioSource LLC ("UBC"), the current operator of the Acthar Hub, and other designated operators of the Program to perform a preliminary assessment of benefit verification for this patient and furnish information requested by the patient's insurer that is available on this form. I understand that insurance verification is ultimately the responsibility of the provider and that third-party reimbursement is affected by a variety of factors. While UBC tries to provide accurate information, they and Mallinckrodt make no representations or warranties as to the accuracy of the information provided.

I understand that representatives from the Program or UBC may contact me or my patient for additional information relating to this prescription. I acknowledge and agree that the designated Specialty Pharmacy receive this prescription via a designated third party, the Program, and that no additional confirmation of receipt of prescription is required by the designated Specialty Pharmacy.

HCP Prescriber Signature - Please sign ONE LINE below

DISPENSE AS WRITTEN _____ DATE _____ **OR** SUBSTITUTIONS ALLOWED _____ DATE _____

Prescriber signature required for consent and to validate prescriptions. Prescriber attests that this is her/his signature. NO STAMPS. By signing, prescriber certifies that the above is medically necessary.



6. DIAGNOSIS AND MEDICAL INFORMATION

Diagnosis Codes

Please check the diagnosis code that corresponds with the patient's condition. Below is a list of common ICD-10 codes and you may also write the patient's diagnosis in the "OTHER" section.

PULMONOLOGY

- SARCOIDOSIS
D86
- SARCOIDOSIS OF LUNG
D86.0
- SARCOIDOSIS OF LYMPH NODES
D86.1
- SARCOIDOSIS OF LUNG WITH SARCOIDOSIS OF LYMPH NODES
D86.2
- SARCOIDOSIS OF SKIN
D86.3
- SARCOIDOSIS OF OTHER SITES
D86.8
- SARCOID MENINGITIS
D86.81

- MULTIPLE CRANIAL NERVE PALSIES IN SARCOIDOSIS
D86.82
- SARCOID IRIDOCYCLITIS
D86.83
- SARCOID PYELONEPHRITIS
D86.84
- SARCOID MYOCARDITIS
D86.85
- SARCOID ARTHROPATHY
D86.86
- SARCOID MYOSITIS
D86.87
- SARCOIDOSIS OF OTHER SITES
D86.89
- SARCOIDOSIS, UNSPECIFIED
D86.9

RHEUMATOLOGY

- ARTHROPATHIC PSORIASIS, UNSPECIFIED
L40.50
- OTHER PSORIATIC ARTHROPATHY
L40.59

- RHEUMATOID ARTHRITIS WITH RHEUMATOID FACTOR OF MULTIPLE SITES WITHOUT ORGAN OR SYSTEMS INVOLVEMENT
M05.79
- RHEUMATOID ARTHRITIS, UNSPECIFIED
M06.9
- SYSTEMIC LUPUS ERYTHEMATOSUS, ORGAN OR SYSTEM INVOLVEMENT UNSPECIFIED
M32.10
- GLOMERULAR DISEASE IN SYSTEMIC LUPUS ERYTHEMATOSUS
M32.14
- SYSTEMIC LUPUS ERYTHEMATOSUS, UNSPECIFIED
M32.9
- OTHER DERMATOMYOSITIS WITH MYOPATHY
M33.12
- POLYMYOSITIS, ORGAN INVOLVEMENT UNSPECIFIED
M33.20
- POLYMYOSITIS WITH MYOPATHY
M33.22

OPHTHALMOLOGY

- NEUROMYELITIS OPTICA [DEVIC]
G36.0
- UNSPECIFIED SCLERITIS, UNSPECIFIED EYE
H15.009
- SCLERITIS WITH CORNEAL INVOLVEMENT, RIGHT EYE
H15.041
- UNSPECIFIED SUPERFICIAL KERATITIS, BILATERAL
H16.103
- FILAMENTARY KERATITIS, BILATERAL
H16.123
- PUNCTATE KERATITIS, RIGHT EYE
H16.141
- PUNCTATE KERATITIS, LEFT EYE
H16.142
- PUNCTATE KERATITIS, BILATERAL
H16.143
- OTHER KERATOCONJUNCTIVITIS, BILATERAL
H16.293
- UNSPECIFIED INTERSTITIAL KERATITIS, RIGHT EYE
H16.301

- OTHER KERATITIS
H16.8
- PRIMARY IRIDOCYCLITIS, LEFT EYE
H20.012
- RECURRENT ACUTE IRIDOCYCLITIS, LEFT EYE
H20.022
- SECONDARY NONINFECTIOUS IRIDOCYCLITIS, RIGHT EYE
H20.041
- CHRONIC IRIDOCYCLITIS, RIGHT EYE
H20.11
- CHRONIC IRIDOCYCLITIS, LEFT EYE
H20.12
- CHRONIC IRIDOCYCLITIS, BILATERAL
H20.13
- UNSPECIFIED IRIDOCYCLITIS
H20.9
- UNSPECIFIED CHORIORETINAL INFLAMMATION, BILATERAL
H30.93
- RETINAL VASCULITIS, BILATERAL
H35.063

- PANUVEITIS, RIGHT EYE
H44.111
- PANUVEITIS, LEFT EYE
H44.112
- PANUVEITIS, BILATERAL
H44.113
- SYMPATHETIC UVEITIS, UNSPECIFIED EYE
H44.139
- RETROBULBAR NEURITIS, RIGHT EYE
H46.11
- RETROBULBAR NEURITIS, LEFT EYE
H46.12
- OTHER OPTIC NEURITIS
H46.8
- UNSPECIFIED OPTIC NEURITIS
H46.9
- OTHER IRREGULAR EYE MOVEMENTS
H55.89

Other: _____

ORGAN INVOLVEMENT

- LUNGS
- LYMPH NODES

- SKIN AND TISSUES
- EYES
- HEART

- BRAIN AND NERVOUS SYSTEM
- BONES, JOINTS, CARTILAGE, LIGAMENTS, TENDONS AND MUSCLES

- SPLEEN
- LIVER
- KIDNEYS AND URINARY TRACT

- SALIVARY GLANDS
- SINUSES

OTHER: _____

7. HISTORY OF CORTICOSTEROID USE (IF APPLICABLE) PLEASE ADD DETAILS IN SECTION 9 BELOW.

Please check all that apply:

- A corticosteroid **was** tried with the following response(s):
- Corticosteroid use failed, but same response not expected with Acthar
- Patient hypersensitive or allergic to corticosteroids
- Patient intolerant of corticosteroids
- Other: _____

OR

- A corticosteroid **was not** tried due to the following response(s):
- Corticosteroid use is contraindicated for this patient
- Intravenous access is not possible for this patient
- Patient has known intolerance to corticosteroids
- Other: _____

8. CONCURRENT MEDICATIONS

9. RELEVANT TREATMENT HISTORY (INCLUDING RECENT STEROID HISTORY. ATTACH ADDITIONAL PAGES AS NECESSARY.)

Therapy Name	Dose	Start Date	Stop Date (if applicable)	Explain Outcome With Detail (eg, type of outcome)

OTHER RELEVANT CLINICAL INFORMATION

HCP SIGNATURE: REQUIRED FOR DOCUMENTATION

I verify that the patient and healthcare provider information on this enrollment form was completed by me or at my direction and that the information contained herein is complete and accurate to the best of my knowledge. I certify that my patient has agreed in writing to be contacted by Program administrators or UBC and be furnished with Program or other information or materials.

NAME _____ SIGNATURE _____ DATE _____



Patient Name: _____ **Date of Birth:** _____

10. PATIENT AUTHORIZATION(S)

Patient Consent to allow Acthar Support Team to work together with your insurance provider, pharmacy, advocacy organization and others to provide support on your behalf.

By signing this authorization, I authorize my physician(s), my health insurance company and my pharmacy providers (collectively, "Designated Parties") to disclose to Mallinckrodt ARD Inc. ("Mallinckrodt"), the distributor of Acthar, and its agents, authorized designees and contractors, including Mallinckrodt reimbursement support personnel and United BioSource LLC ("UBC") or any other operator of the Acthar Hub on behalf of Mallinckrodt (collectively, "Manufacturer Parties"), health information relating to my medical condition, treatment and insurance coverage (my "Health Information") in order for them to (1) provide certain services to me, including reimbursement and coverage support, patient assistance and access programs, medication shipment tracking, and home injection training, (2) provide me with support services and information associated with my Acthar therapy, (3) serve internal business purposes, such as marketing research, internal financial reporting and operational purposes, and (4) carry out the Manufacturer Parties' respective legal responsibilities.

Once my Health Information has been disclosed to Manufacturer Parties, I understand that it may be redisclosed by them and no longer protected by federal and state privacy laws. However, Manufacturer Parties agree to protect my Health Information by using and disclosing it only for the purposes detailed in this authorization or as permitted or required by law.

I understand that I may refuse to sign this authorization and that my physician and pharmacy will not condition my treatment on my agreement to sign this authorization form, and my health plan or health insurance company will not condition payment for my treatment, insurance enrollment or eligibility for insurance benefits on my agreement to sign this authorization form. I understand that my pharmacies and other Designated Parties may receive payment in connection with the disclosure of my Health Information as provided in this authorization. I understand that I am entitled to receive a copy of this authorization after I sign it.

I may revoke (withdraw) this authorization at any time by mailing a letter to the Acthar Hub, 255 Technology Park, Lake Mary, FL 32746. Revoking this authorization will end further disclosure of my Health Information to Manufacturer Parties by my pharmacy, physicians, and health insurance company when they receive a copy of the revocation, but it will not apply to information they have already disclosed to Manufacturer Parties based on this authorization. I also know I may cancel my enrollment in a patient support program at any time in writing by contacting Mallinckrodt via fax at 1-877-937-2284 or by calling the Acthar Hub at 1-888-435-2284. This authorization is in effect for 5 years unless a shorter period is provided for by state law or until the conclusion of any ongoing coverage support, whichever is longer, once I have signed it unless I cancel it before then.

THIS SECTION MUST BE COMPLETED IN ITS ENTIRETY, INCLUDING DATE

▶ _____
PATIENT NAME OR LEGAL REPRESENTATIVE PATIENT SIGNATURE IF LEGAL REPRESENTATIVE, RELATIONSHIP TO PATIENT DATE

Patient Consent to receive additional information from Mallinckrodt such as education on your disease and Acthar.

I authorize Mallinckrodt and its partners to use, disclose, and/or transfer the personal information I supply (1) to contact me and provide me with informational and marketing materials and clinical trial opportunities related to my condition or treatment by any means of communication, including but not limited to text, email, mail, or telephone; (2) to help Mallinckrodt improve, develop, and evaluate products, services, materials, and programs related to my condition or treatment; (3) to enroll me in and provide me with Acthar-related programs and services that I may select or refuse at any time; (4) to disclose my enrollment and use of these services to my healthcare providers and insurers; and (5) to use my information that cannot identify me for scientific and market research. This authorization will remain in effect until I cancel it, which I may do at any time in writing by contacting Mallinckrodt via fax at 1-877-937-2284 or by calling the Acthar Hub at 1-888-435-2284. I may request a copy of this signed authorization.

▶ _____
PATIENT NAME OR LEGAL REPRESENTATIVE PATIENT SIGNATURE IF LEGAL REPRESENTATIVE, RELATIONSHIP TO PATIENT DATE

If patient is not present to sign the form, send them to
ActharConsent.com
and have them sign electronically.

RESOURCE PAGE. DO NOT NEED TO FAX BACK.

RHEUMATOLOGY,
continued

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED HIP
M08.059

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, RIGHT KNEE
M08.061

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, LEFT KNEE
M08.062

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED KNEE
M08.069

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, RIGHT ANKLE AND FOOT
M08.071

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, LEFT ANKLE AND FOOT
M08.072

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED ANKLE AND FOOT
M08.079

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, VERTEBRAE
M08.08

UNSPECIFIED JUVENILE RHEUMATOID ARTHRITIS, MULTIPLE SITES
M08.09

JUVENILE ANKYLOSING SPONDYLITIS
M08.1

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED SITE
M08.20

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, RIGHT SHOULDER
M08.211

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT SHOULDER
M08.212

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED SHOULDER
M08.219

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, MULTIPLE SITES
M08.29

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT ELBOW
M08.221

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT ELBOW
M08.222

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED ELBOW
M08.229

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, RIGHT WRIST
M08.231

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT WRIST
M08.232

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED WRIST
M08.239

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED WRIST
M08.239

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, RIGHT HAND
M08.241

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT HAND
M08.242

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED HAND
M08.249

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, RIGHT HIP
M08.251

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT HIP
M08.252

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED HIP
M08.259

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, RIGHT KNEE
M08.261

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT KNEE
M08.262

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED KNEE
M08.269

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED KNEE
M08.269

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, RIGHT ANKLE AND FOOT
M08.271

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, LEFT ANKLE AND FOOT
M08.272

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, UNSPECIFIED ANKLE AND FOOT
M08.279

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, VERTEBRAE
M08.28

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, MULTIPLE SITES
M08.29

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, VERTEBRAE
M08.28

JUVENILE RHEUMATOID ARTHRITIS WITH SYSTEMIC ONSET, MULTIPLE SITES
M08.29

JUVENILE RHEUMATOID POLYARTHRITIS (SERONEGATIVE)
M08.3

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED SITE
M08.40

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, RIGHT SHOULDER
M08.411

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, LEFT SHOULDER
M08.412

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED SHOULDER
M08.419

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, RIGHT ELBOW
M08.421

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, LEFT ELBOW
M08.422

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED ELBOW
M08.429

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, RIGHT WRIST
M08.431

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, LEFT WRIST
M08.432

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED WRIST
M08.439

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, RIGHT HAND
M08.441

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, LEFT HAND
M08.442

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED HAND
M08.449

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, RIGHT HIP
M08.451

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, LEFT HIP
M08.452

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED HIP
M08.459

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, RIGHT KNEE
M08.461

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, LEFT KNEE
M08.462

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED KNEE
M08.469

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, RIGHT ANKLE AND FOOT
M08.471

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, LEFT ANKLE AND FOOT
M08.472

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED ANKLE AND FOOT
M08.479

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, UNSPECIFIED ANKLE AND FOOT
M08.479

PAUCIARTICULAR JUVENILE RHEUMATOID ARTHRITIS, VERTEBRAE
M08.48

SYSTEMIC LUPUS ERYTHEMATOSUS, ORGAN OR SYSTEM INVOLVED UNSPECIFIED
M32.10

ENDOCARDITIS IN SYSTEMIC LUPUS ERYTHEMATOSUS
M32.11

PERICARDITIS IN SYSTEMIC LUPUS ERYTHEMATOSUS
M32.12

LUNG INVOLVEMENT IN SYSTEMIC LUPUS ERYTHEMATOSUS
M32.13

GLOMERULAR DISEASE IN SYSTEMIC LUPUS ERYTHEMATOSUS
M32.14

TUBULO-INTERSTITIAL NEUROPATHY IN SYSTEMIC LUPUS ERYTHEMATOSUS
M32.15

OTHER ORGAN OR SYSTEM INVOLVED IN SYSTEMIC LUPUS ERYTHEMATOSUS
M32.19

OTHER FORMS OF SYSTEMIC LUPUS ERYTHEMATOSUS
M32.8

SYSTEMIC LUPUS ERYTHEMATOSUS, UNSPECIFIED
M32.9

JUVENILE DERMATOPOLYMYOSITIS, ORGAN INVOLVEMENT UNSPECIFIED
M33.00

JUVENILE DERMATOPOLYMYOSITIS WITH RESPIRATORY INVOLVEMENT
M33.01

JUVENILE DERMATOPOLYMYOSITIS WITH MYOPATHY
M33.02

JUVENILE DERMATOPOLYMYOSITIS WITH OTHER ORGAN INVOLVEMENT
M33.09

OTHER DERMATOPOLYMYOSITIS, ORGAN INVOLVEMENT UNSPECIFIED
M33.10

OTHER DERMATOPOLYMYOSITIS WITH RESPIRATORY INVOLVEMENT
M33.11

OTHER DERMATOPOLYMYOSITIS WITH MYOPATHY
M33.12

OTHER DERMATOPOLYMYOSITIS WITH OTHER ORGAN INVOLVEMENT
M33.19

POLYMYOSITIS, ORGAN INVOLVEMENT UNSPECIFIED
M33.20

POLYMYOSITIS WITH RESPIRATORY INVOLVEMENT
M33.21

POLYMYOSITIS WITH MYOPATHY
M33.22

POLYMYOSITIS WITH OTHER ORGAN INVOLVEMENT
M33.29

DERMATOPOLYMYOSITIS, UNSPECIFIED, ORGAN INVOLVEMENT UNSPECIFIED
M33.90

DERMATOPOLYMYOSITIS, UNSPECIFIED WITH RESPIRATORY INVOLVEMENT
M33.91

DERMATOPOLYMYOSITIS, UNSPECIFIED WITH MYOPATHY
M33.92

DERMATOPOLYMYOSITIS, UNSPECIFIED WITH OTHER ORGAN INVOLVEMENT
M33.99

DERMATO(POLY)MYOSITIS IN NEOPLASTIC DISEASE
M36.0

ANKYLOSING SPONDYLITIS OF MULTIPLE SITES IN SPINE
M45.0

ANKYLOSING SPONDYLITIS OF OCCIPITO-ATLANTO-AXIAL REGION
M45.1

ANKYLOSING SPONDYLITIS OF CERVICAL REGION
M45.2

ANKYLOSING SPONDYLITIS OF CERVICOTHORACIC REGION
M45.3

ANKYLOSING SPONDYLITIS OF THORACIC REGION
M45.4

ANKYLOSING SPONDYLITIS OF THORACOLUMBAR REGION
M45.5

ANKYLOSING SPONDYLITIS OF LUMBAR REGION
M45.6

ANKYLOSING SPONDYLITIS OF LUMBOSACRAL REGION
M45.7

ANKYLOSING SPONDYLITIS SACRAL AND SACROCOCCYGEAL REGION
M45.8

ANKYLOSING SPONDYLITIS OF UNSPECIFIED SITES IN SPINE
M45.9

OTHER SERUM REACTION DUE TO OTHER SERUM, INITIAL ENCOUNTER
T80.69XA

OPHTHALMOLOGY

SARCOID IRIDOCYCLITIS
D86.83

DISCOID LUPUS ERYTHEMATOSUS OF RIGHT UPPER EYELID
H01.121

DISCOID LUPUS ERYTHEMATOSUS OF RIGHT LOWER EYELID
H01.122

DISCOID LUPUS ERYTHEMATOSUS OF RIGHT EYE, UNSPECIFIED EYELID
H01.123

DISCOID LUPUS ERYTHEMATOSUS OF LEFT UPPER EYELID
H01.124

DISCOID LUPUS ERYTHEMATOSUS OF LEFT LOWER EYELID
H01.125

DISCOID LUPUS ERYTHEMATOSUS OF LEFT EYE, UNSPECIFIED EYELID
H01.126

DISCOID LUPUS ERYTHEMATOSUS OF UNSPECIFIED EYE, UNSPECIFIED EYELID
H01.129

OTHER SPECIFIED INFLAMMATIONS OF EYELID
H01.8

UNSPECIFIED INFLAMMATION OF EYELID
H01.9

CHRONIC DACRYOADENITIS, RIGHT LACRIMAL GLAND
H04.021

CHRONIC DACRYOADENITIS, LEFT LACRIMAL GLAND
H04.022

CHRONIC DACRYOADENITIS, BILATERAL LACRIMAL GLAND
H04.023

CHRONIC DACRYOCYSTITIS OF LEFT LACRIMAL PASSAGE
H04.411

CHRONIC DACRYOCYSTITIS OF RIGHT LACRIMAL PASSAGE
H04.412

CHRONIC DACRYOCYSTITIS OF UNSPECIFIED LACRIMAL PASSAGE
H04.419

UNSPECIFIED ACUTE INFLAMMATION OF ORBIT
H05.00

TENONITIS OF RIGHT ORBIT
H05.041

TENONITIS OF LEFT ORBIT
H05.042

TENONITIS OF BILATERAL ORBITS
H05.043

TENONITIS OF UNSPECIFIED ORBIT
H05.049

UNSPECIFIED CHRONIC INFLAMMATORY DISORDERS OF ORBIT
H05.10

GRANULOMA OF RIGHT ORBIT
H05.111

GRANULOMA OF LEFT ORBIT
H05.112

GRANULOMA OF BILATERAL ORBITS
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PULMONOLOGY

SARCOIDOSIS OF LUNG
D86.0

SARCOIDOSIS OF LUNG
WITH SARCOIDOSIS OF
LYMPH NODES
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INDICATIONS AND USAGE

- **Infantile spasms:** H.P. Acthar Gel (repository corticotropin injection) is indicated as monotherapy for the treatment of infantile spasms in infants and children under 2 years of age
- **Multiple Sclerosis:** H.P. Acthar Gel is indicated for the treatment of acute exacerbations of multiple sclerosis in adults. Controlled clinical trials have shown H.P. Acthar Gel to be effective in speeding the resolution of acute exacerbations of multiple sclerosis. However, there is no evidence that it affects the ultimate outcome or natural history of the disease
- **Rheumatic Disorders:** As adjunctive therapy for short-term administration (to tide the patient over an acute episode or exacerbation) in: Psoriatic arthritis; Rheumatoid arthritis, including juvenile rheumatoid arthritis (selected cases may require low-dose maintenance therapy), Ankylosing spondylitis
- **Collagen Diseases:** During an exacerbation or as maintenance therapy in selected cases of: systemic lupus erythematosus, systemic dermatomyositis (polymyositis)
- **Dermatologic Diseases:** Severe erythema multiforme, Stevens-Johnson syndrome
- **Allergic States:** Serum sickness
- **Ophthalmic Diseases:** Severe acute and chronic allergic and inflammatory processes involving the eye and its adnexa such as: keratitis; iritis, iridocyclitis, diffuse posterior uveitis and choroiditis, optic neuritis, chorioretinitis; anterior segment inflammation
- **Respiratory Diseases:** Symptomatic sarcoidosis
- **Edematous State:** To induce a diuresis or a remission of proteinuria in the nephrotic syndrome without uremia of the idiopathic type or that due to lupus erythematosus

IMPORTANT SAFETY INFORMATION

Contraindications

- Acthar should never be administered intravenously
- Administration of live or live attenuated vaccines is contraindicated in patients receiving immunosuppressive doses of Acthar
- Acthar is contraindicated where congenital infections are suspected in infants
- Acthar is contraindicated in patients with scleroderma, osteoporosis, systemic fungal infections, ocular herpes simplex, recent surgery, history of or the presence of a peptic ulcer, congestive heart failure, uncontrolled hypertension, primary adrenocortical insufficiency, adrenocortical hyperfunction or sensitivity to proteins of porcine origins

Warnings and Precautions

- The adverse effects of Acthar are related primarily to its steroidogenic effects
- Acthar may increase susceptibility to new infection or reactivation of latent infections
- Suppression of the hypothalamic-pituitary-axis (HPA) may occur following prolonged therapy with the potential for adrenal insufficiency after withdrawal of the medication. Adrenal insufficiency may be minimized by tapering of the dose when discontinuing treatment. During recovery of the adrenal gland patients should be protected from the stress (e.g. trauma or surgery) by the use of corticosteroids. Monitor patients for effects of HPA suppression after stopping treatment
- Cushing's syndrome may occur during therapy but generally resolves after therapy is stopped. Monitor patients for signs and symptoms
- Acthar can cause elevation of blood pressure, salt and water retention, and hypokalemia. Blood pressure, sodium and potassium levels may need to be monitored
- Acthar often acts by masking symptoms of other diseases/disorders. Monitor patients carefully during and for a period following discontinuation of therapy
- Acthar can cause GI bleeding and gastric ulcer. There is also an increased risk for perforation in patients with certain gastrointestinal disorders. Monitor for signs of bleeding
- Acthar may be associated with central nervous system effects ranging from euphoria, insomnia, irritability, mood swings, personality changes, and severe depression, and psychosis. Existing conditions may be aggravated
- Patients with comorbid disease may have that disease worsened. Caution should be used when prescribing Acthar in patients with diabetes and myasthenia gravis
- Prolonged use of Acthar may produce cataracts, glaucoma and secondary ocular infections. Monitor for signs and symptoms
- Acthar is immunogenic and prolonged administration of Acthar may increase the risk of hypersensitivity reactions. Neutralizing antibodies with chronic administration may lead to loss of endogenous ACTH activity
- There is an enhanced effect in patients with hypothyroidism and in those with cirrhosis of the liver
- Long-term use may have negative effects on growth and physical development in children. Monitor pediatric patients
- Decrease in bone density may occur. Bone density should be monitored for patients on long-term therapy
- Pregnancy Class C: Acthar has been shown to have an embryocidal effect and should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus

Adverse Reactions

- Common adverse reactions for Acthar are similar to those of corticosteroids and include fluid retention, alteration in glucose tolerance, elevation in blood pressure, behavioral and mood changes, increased appetite and weight gain
- Specific adverse reactions reported in IS clinical trials in infants and children under 2 years of age included: infection, hypertension, irritability, Cushingoid symptoms, constipation, diarrhea, vomiting, pyrexia, weight gain, increased appetite, decreased appetite, nasal congestion, acne, rash, and cardiac hypertrophy. Convulsions were also reported, but these may actually be occurring because some IS patients progress to other forms of seizures and IS sometimes mask other seizures, which become visible once the clinical spasms from IS resolve

Other adverse events reported are included in the full Prescribing Information.

Please see accompanying full Prescribing Information and Medication Guide.