

D-Penamine[®] Prior Authorization Request Form (Page 1 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information <small>(required)</small>					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information <small>(required)</small>					
Select the diagnosis below: <input type="checkbox"/> Cystinuria <input type="checkbox"/> Severe Active Rheumatoid Arthritis <input type="checkbox"/> Wilson's Disease <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Select if the requested medication is prescribed by or in consultation with one of the following specialists: <input type="checkbox"/> Gastroenterologist <input type="checkbox"/> Nephrologist <input type="checkbox"/> Hepatologist <input type="checkbox"/> Rheumatologist <input type="checkbox"/> Metabolic geneticist					
For Cystinuria, also answer the following: Does the patient have urinary cystine excretion of > 300mg/day? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient had a 30 day trial and failure of potassium citrate or other urinary alkalinizing agent along with sodium and protein-restricted diet and hyperdiuresis (urine output of at least 3L/day)? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the dosing exceed 4gm/day? <input type="checkbox"/> Yes <input type="checkbox"/> No Reauthorization: If this is a reauthorization request, answer the following: Does the patient have urinary cystine excretion of < 200mg/day? [‡] <input type="checkbox"/> Yes <input type="checkbox"/> No [‡] Labs must be submitted for documentation Does the dosing exceed 4gm/day? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For Severe Active Rheumatoid Arthritis, also answer the following: Select if the patient has had <u>30 days</u> trial and failure of the following: <input type="checkbox"/> Cimzia <input type="checkbox"/> Humira <input type="checkbox"/> Simponi Does the dosing exceed 250mg/day for the first month and 1.5gm/day for maintenance therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Reauthorization: If this is a reauthorization request, answer the following: Has the patient had shown a clinically significant improvement in rheumatoid arthritis symptoms with chart notes documenting improvement in symptoms? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the dosing exceed 1.5gm/day for maintenance therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					

D-Penamine[®] Prior Authorization Request Form (Page 2 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

For Wilson's Disease, also answer the following:

Select if the patient's diagnosis of Wilson's disease is confirmed by one of the following*:

- 24-hour urine copper > 40 mcg
- Liver biopsy with copper dry weight > 250 mcg/g
- Presence of Kayser-Fleisher rings
- Serum ceruloplasmin (CPN) < 20mg/dL

*Submit labs or chart notes for documentation

Does the dosing exceed 1.5gm/day? Yes No

Reauthorization:

If this is a reauthorization request, answer the following:

Is the patient's serum copper level < 10 mcg free copper/dL of serum?* Yes No

*Labs must be submitted for documentation

Does the dosing exceed 1.5gm/day? Yes No

Quantity limit requests:

What is the quantity requested per DAY? _____

Previous therapies failed and/or therapies currently used in combination with the requested medication (*List ALL medications tried or authorization process will be delayed*): _____

Is the prescribed dose higher than the maximum dose recommendation in FDA-approved labeling (i.e., the package insert)? Yes No

If **Yes**, please provide documentation to support the safety and efficacy of the higher dose (such as evidence from practice guidelines or clinical trials from peer-reviewed medical literature).

** Please note: Chart documentation of the above is required to be submitted along with this fax

****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.

Prescriber Signature: _____ Date: _____

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.