



## PRIOR AUTHORIZATION REQUEST

### Strensiq

#### Patient Information:

Name:	
Member ID:	
Address:	
City, State, Zip:	
Date of Birth:	

#### Prescriber Information:

Name:	
NPI:	
Phone Number:	
Fax Number:	
Address:	
City, State, Zip:	

#### Requested Medication

Rx Name:	
Rx Strength:	
Rx Quantity:	
Rx Frequency:	
Rx Route of Administration:	
Diagnosis and ICD Code:	

Your patient's prescription benefit requires that we review certain requests for coverage with the prescriber. You have prescribed a medication for your patient that requires Prior Authorization before benefit coverage or coverage of additional quantities can be provided. Please complete the following questions then fax this form to the toll-free number listed below. Upon receipt of the completed form, prescription benefit coverage will be determined based on the plan's rules.

**SECTION A:** Please note that supporting clinical documentation is required for **ALL PA** requests. Pharmacy prior authorization reviews can be subject to trial with additional medications that are not listed within the criteria. The policies are subject to change based on COMAR requirements, MDH transmittals and updates to treatment guidelines.

- |   |  |             |
|---|--|-------------|
| 1 | What is the diagnosis or indication?<br><input type="checkbox"/> Perinatal/infantile-onset hypophosphatasia (If checked, go to 2)<br><br><input type="checkbox"/> Juvenile-onset hypophosphatasia (If checked, go to 2)<br><br><input type="checkbox"/> Other (If checked, no further questions) |             |
| 2 | Did the patient have a disease onset at 18 years of age or younger?<br>[If no, no further questions.]  | Yes      No |

If you have any  
questions, call:  
1-888-258-8250

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- |   |  |     |    |
|---|--|-----|----|
| 3 | Has the requested medication been prescribed by or in consultation with a geneticist, endocrinologist, a metabolic disorder sub-specialist, or a physician who specializes in the treatment of hypophosphatasia or related disorders?<br>[If no, no further questions.]  | Yes | No |
| 4 | Has the patient undergone molecular genetic testing which documents a tissue non-specific alkaline phosphatase (ALPL) gene mutation?<br>[If yes, skip to question 7.]  | Yes | No |
| 5 | Does the patient have low baseline serum alkaline phosphatase activity?<br>[If yes, skip to question 7.]   | Yes | No |
| 6 | Does the patient have an elevated level of a tissue non-specific alkaline phosphatase substrate (i.e., serum pyridoxal 5'-phosphate, serum or urinary inorganic pyrophosphate, urinary phosphoethanolamine)?<br>[If no, no further questions.]   | Yes | No |
| 7 | Does the patient have a history of or currently have clinical manifestations consistent with hypophosphatasia (for example, skeletal abnormalities, premature tooth loss, muscle weakness, poor feeding, failure to thrive, respiratory problems, Vitamin B6-dependant seizures)?<br>[If yes, no further questions.] | Yes | No |
| 8 | Does the patient have a family history (parent or sibling) of hypophosphatasia without current clinical manifestations of hypophosphatasia?  | Yes | No |

**Please document the diagnoses, symptoms, and/or any other information important to this review:**

### SECTION B: Physician Signature

PHYSICIAN SIGNATURE

DATE

**FAX COMPLETED FORM TO: 1-833-896-0656**

**Disclaimer:** An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior authorization as per Plan policy and procedures.

**Confidentiality:** The information contained in this transmission is confidential and may be protected under

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questions, call:  
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