OPTIMA HEALTH PLAN

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions:</u> The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information (<u>including phone and fax #s</u>) on this form is correct. <u>If information provided is not complete, correct, or legible, authorization may be delayed.</u>

<u>Drug Requested</u>: **Endari**[™] (L-glutamine oral powder)

Drug	Information: Authorization may be delayed if incomplete.		
Drug 1	Form/Strength:		
Dosing Schedule: Length of Therapy:			
Diagn	osis: ICD Code, if applicable:		
each li	ICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support ne checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided est may be denied.		
<u>Initia</u>	<u>l Approval</u> – 6 months		
	Member must be 5 years of age or older. Please note member's current weight:		
	AND		
	Member will be dosed as follows (dose above maximum recommended for weight will not be approved):		
	□ <30 kg: 5 g (1 packet) twice daily (total dose 10 g/day)		
	□ 30 to 65 kg: 10 g (2 packets) twice daily (total dose 20 g/day)		
	□ >65 kg: 15 g (3 packets) twice daily (total dose 30 g/day)		
	Member must have a diagnosis of sickle cell disease		
	AND		
	Provider must be a hematologist or oncologist specializing in treatment of sickle cell disease		
	AND		
	Member must have been compliant with hydroxyurea for the last 90 days (compliance will be verified by pharmacy paid claims)		
	AND		
	Member has experienced at least 2 documented sickle cell crises (SCC) events within the preceding 12 months		
	AND		

(Continued on next page)

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Medical chart notes from the last 12 months must be submitted for documentation of frequency of SCC
events and emergency department or other medical facility visits due to SCC events

AND

☐ Member will not take Endari[™] concomitantly with Oxbryta[®] (voxelotor) tablets, Adakveo[®] (crizanlizumab) infusions, or any experimental treatment for sickle cell disease complications

<u>Continuation of Therapy Approval</u> – **12 months**. Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

☐ All of the initial authorization criteria for Endari continues to be met

AND

□ Patient must have been compliant with <u>BOTH</u> Endari[™] <u>AND</u> hydroxyurea since last approval (monthly pharmacy claims must be noted)

AND

□ The frequency of the member's sickle cell crisis events must have decreased since last approval of Endari[™] **OR** have been maintained below the number of events at baseline (medical chart notes must be submitted to document frequency of SCC events and emergency department or other medical facility visits due to SCC events since last approval of Endari[™])

Not all drugs may be covered under every Plan

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

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.

Patient Name:	
Member Optima #:	
Prescriber Name:	
Prescriber Signature:	
Office Contact Name:	
Phone Number:	Fax Number:
DEA OD NDI #.	

*Approved by Pharmacy and Therapeutics Committee: 41/16/2018; 3/21/2019; 3/192020

REVISED/UPDATED: 2/22/2018; 5/10/2019; 6/11/2020