HEALTHY U MEDICAID

PRIOR AUTHORIZATION REQUEST FORM

HEREDITARY ANGIOEDEMA AGENTS

Berinert®, Cinryze®, icatibant, Firazyr®, Haegarda®, Kalbitor®, Takhzyro®

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.									
If you have prior authorization questions, please call for assistance: 385-425-5094									
Disclaimer: Prior Authorization request forms are subject to change in accordance with Federal and State notice requirements.									
		T.,		T					
Date:		Member Name:		ID#:					
DOB:		Gender:		Physician:					
Office Phone:		Office Fax:		Office Contact:					
Height/Weight:									
Preferred: ☐ Berinert®(C1 esterase inhibitor [human])*, ☐ Haegarda® (C1 esterase inhibitor [human]), ☐ icatibant Non-preferred: ☐ Cinryze® (C1 esterase inhibitor subcutaneous [human]) ☐ Firazyr® (icatibant), ☐ Kalbitor® (ecallantide), ☐ Takhzyro® (lanadelumab) *preferred for specified populations. Refer to medication use policy. Dosing/Frequency: ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐									
If the request is for reauthorization, proceed to reauthorization section									
	Question	ns	Yes	No	Comments/Notes				
1.	Is the request for treatment of He	reditary Angioedema (HAE)?							
2.	Is the requesting provider a board allergist?	-certified immunologist or							
3.	Does the member have clinical pro HAE subtype (HAE I, HAE II, or HAI confirmed by repeat blood testing	E with normal C1INH)			Please provide documentation				
4.	Has the member's diagnosis of He confirmed with complement 4 (C4 levels?				Please provide documentation				
5.	Has the member had a trial and fa antihistamines, glucocorticoids, a	_			Please provide documentation				
6.	Is the member currently taking AC containing oral medications?								
7.	Has the member's attack frequent documented?	cy, severity, and location been			Please provide documentation				

8.	Is the member/caregiver able and ready to administer medication at home?							
9.	For acute HAE attack treatment: Does the member have a history of at least one attack per year?			Please provide documentation				
10.	For long-term prophylaxis of HAE attacks: Does the member have a history of two acute severe attacks per month or at least 5 attacks of moderate severity per month on average?			Please provide documentation				
11.	For long-term prophylaxis of HAE attacks: Has the member tried and failed, or have a contraindication to, danazol therapy?			Please provide documentation				
12.	For long-term prophylaxis of HAE attacks: Does laboratory test show the member has not experienced HAE attacks due to preventable triggers, such as helicobacter pylori infections in members with gastrointestinal attacks?			Please provide documentation				
REAUTHORIZATION								
1.	Is the request for reauthorization of therapy?							
2.	Has the member experienced unacceptable toxicity (e.g. hypersensitivity reactions, serious thrombotic events, significantly elevated hepatic serum transaminases) to the drug?							
3.	For acute HAE attack treatment: Does documentation show that the member continues to experience at least one acute HAE attack per year AND is the request for a refill due to a documented attack OR has the medication on hand reached the expiration date?			Please provide documentation				
4.	For long-term prophylaxis of HAE attacks: Has the provider evaluated the member's need for long-term prophylaxis at least once per year?							
5.	For long-term prophylaxis of HAE attacks: Has the member had significant improvements in severity and duration of attacks compared to baseline?			Please provide documentation				
What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.								
Additional information:								
Physician's Signature:								

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Policy PHARM-HU-031 Origination Date: 01/01/2022 Reviewed/Revised Date: 11/08/2023 Next Review Date: 11/08/2024 Current Effective Date: 12/01/2023

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