Vyndaqel (tafamidis) **Prior Authorization Request Form** Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

EMBER'S FIRST NAME:			
etely and legibly. Attach any additional documentation that support the authorization request). Information contained			
FIRST NAME:			
DATE OF BIRTH:			
STATE: ZIP CODE:			
WEIGHT (LB/KG):			
submit a PHI Disclosure Authorization form with this rapeutics.com/NOPP			
ICABLE):			
FIRST NAME:			
EMAIL ADDRESS:			
DEA NUMBER:			
FAX NUMBER:			
STATE: ZIP CODE:			
OFFICE CONTACT PERSON:			

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MEMBER'S LAST NAME: MEMBER'S FIRST NAME:						
MEDICATION OR MEDICAL DISPENSING INFORMATION						
MEDICATION NAME:						
DOSE/STRENGTH:	FREQU	ENCY:	LENGTH OF THERAPY/REFIL	LLS:	QUANTITY:	
□ NEW THERAPY □ RENEWAL IF RENEWAL, DATE THERAPY INITIATED:						
DURATION OF THERAPY		<u> </u>				
1. HAS THE PATIENT T YES (IF YES, COMP			ICATIONS FOR	THIS CC	ONDITION?	
Medication/Therapy (Spec Drug Name And Dosage):			Response/Reason For Failure/Allergy:			
2. LIST DIAGNOSES:				ICD-10	:	
☐ Transthyretin amyloid	cardiomy	opathy(wild-type	or hereditary)			
Other diagnosis:						
ICD-10 CODE(S):						
3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.						
Clinical Information:						
Is patient going to be using drug in combination with a clinical trial? Yes No						
Is patient currently taking A	ttruby(ac	coramidis)? 🗌 Yes	□No			
If Yes to the above question, will the Attruby(acoramdis) be discontinued when Vyndamax/Vyndaqel(tafamidis) is started? Yes No						
Is patient currently taking Amvuttra(vutrisiran)? Yes No						
If Yes to the above question, will the Amvuttra(vutrisiran) be discontinued when Vyndamax/Vyndaqel(tafamidis) is started? Yes No						
Does patient have a history of heart failure with at least one prior hospitalization for heart failure? Yes No						
Does patient have clinical evidence of heart failure without hospitalization(defined as signs and symptoms of volume overload or elevated intracardiac pressures requiring treatment with a diuretic)?						



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Is patient's echocardiogram consistent with or suggestive of amyloidosis? Yes No Please submit echocardiogram report.
Does patient have evidence of a left ventricular wall (interventricular septum or left ventricular posterior wall) thickness ≥12 mm? □ Yes □ No Please submit documentation.
Does patient have an N-terminal pro-B-type natriuretic peptide(NT-proBNP) level greater than or equal to 600pg/mL? Yes No Please submit lab report.
Does patient have a B-type natriuretic peptide(BNP) level greater than or equal to 100pg/ml? Yes No Please submit lab report.
If patient had used Attruby(acoramidis), prior to requesting Vyndamax/Vyndaqel, was initial pre-Attruby(acoramidis) NT-proBNP greater than or equal to 600 pg/mL ? ☐ Yes ☐ No Please submit lab report.
Does patient have a New York Heart Association(NYHA) class I, II or III disease? Yes No
Does patient have a confirmed transthyretin precursor protein present via a Grade 2 or Grade 3 positive Tc-pyrophosphate(PYP) scan? Yes No Please submit imaging report.
Are patient's serum immunofixation results within normal range? Yes No Please submit lab report.
Are patient's serum immunofixation results above the upper range of normal listed on the lab report? Yes No Please submit lab report.
Are patient's urine immunofixation results within normal range? Yes No Please submit lab report.
Are patient's urine immunofixation results above the upper range of normal listed on the lab report? Yes No Please submit lab report.
Are patient's serum electrophoresis/free light-chain assay results within normal range? 🗌 Yes 🔝 No
Please submit lab report.
Are patient's serum electrophoresis/free light-chain assay results above the upper range of normal listed on the lab report? Yes No Please submit lab report.
Is patient's free light-chain level within normal range? Yes No Please submit lab report.
Is patient's free light-chain level above the upper range of normal on the lab report?
Please submit lab report.



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Does patient have a confirmed transthyretin precursor protein present via a Grad pyrophosphate(PYP) scan? Yes No Please submit imaging report.	e 1 positive Tc-
Is patient's ATTR amyloid histologically confirmed and typed from an endomyoca specimen? Yes No Please submit tissue biopsy.	ardial tissue biopsy
Is patient's ATTR amyloid histologically confirmed and typed from ANY tissue biopsy.	opsy specimen?
Does a hematology consultation report rule out light-chain disease? Yes report.	No <i>Please submit</i>
Are there any other comments, diagnoses, symptoms, medications tried or failed information the physician feels is important to this review?	, and/or any other
Please note: Not all drugs/diagnosis are covered on all plans. This request may b required information is received.	e denied unless all
ATTESTATION: I attest the information provided is true and accurate to the best understand that the Health Plan, insurer, Medical Group or its designees may per request the medical information necessary to verify the accuracy of the information.	form a routine audit and
Prescriber Signature or Electronic I.D. Verification:	Date:
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FAX THIS FORM TO: 800-424-7640

MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program

Attn: CP-4201 P.O. Box 64811 St. Paul, MN 55164-0811 **Phone**: 877-228-7909

