HYPERLIPIDEMIA AGENTS PRIOR AUTHORIZATION REQUEST

PRESCRIBER FAX FORM

ONLY the prescriber may complete and fax this form. This form is for prospective, concurrent, and retrospective reviews.

Incomplete forms will be returned for additional information. The following documentation is required for prior authorization consideration. For formulary information and to download additional forms, please visit https://www.bcbstx.com/provider/medicaid/pharmacy/rx-prior-auth

PATIENT AND INSURANCE INFO	RMATION				Today	's Date	:	
Patient Name (First):	Last:				M:	DOB	(mm/dd/yy):	
Patient Address:		City, State, Zip:			Pati	Patient Telephone:		
BCBSTX ID Number:	CBSTX ID Number:			Group Number:				
PRESCRIBER/CLINIC INFORMAT	ION							
Prescriber Name:	Prescriber NPI#:		Specialty:			Contact Name:		
Clinic Name:			Clinic Address:					
City, State, Zip:			Phone #:		Sec	Secure Fax #:		
PLEASE ATTACH ANY ADDITION	AL INFOR	MATION THAT S	SHOULD	BE CONSI		H THIS	REQUEST	
Patient's Diagnosis (please check	one of the	following):						
Diagnosis of Heterozygous Far	nilial Hype	rcholesteremia		D	ate of diagno	sis:		
Clinical Atherosclerotic Cardiov	ascular Di	sease		D	ate of diagno	sis:		
Diagnosis of Homozygous Fam	ilial Hyper	cholesteremia		D	ate of diagno	sis:		
Diagnosis of Primary Hyperlipic	lemia			D	ate of diagno	sis:		
Other, please specify ICD code	plus desc	ription				_ Date	of diagnosis:	
Medication Requested:			Strength:					
Dosing Schedule:	Dosing Schedule: Quantity per Month:							
Please indicate PSCK9 Treatment	Status:	🗌 Initial 🛛 🗋 Co	ontinuation	n; Date of t	reatment initi	ation: _		
Expedited/Urgent Review Requ	iested: By	checking this boy	x and sign	ing below,	I certify that a	applying	the standard review time	
frame may seriously jeopardize the		-	-					
Signature of prescriber or presc	riber's de	signee:			Da	te:		
Section 1. Drug Treatment Histo	ry (comple	ete as applicable):	:					
					Start date		End date* (if applicable)	
Drug		Las	t prescrib	ed dose	(mm/dd/ccy	у)	(mm/dd/ccyy or N/A)	
atorvastatin								
ezetimibe								
☐ rosuvastatin								
other (please specify):								
other (please specify):								
other (please specify):								
*For current therapy, indicate "N/A	" for "End o	date".						
Please continue to next page.								

Pat	ient Name (First):	Last:	M:	DOB (mm/dd/yy):		
1.	1. Is the patient currently treated with the requested medication?					
If yes, when was treatment with the requested medication started:						
	If yes, has the patient shown clinical response since initiating therapy?					
2.	2. Is the patient currently pregnant?				🗌 No	
3.	3. Does the patient have a diagnosis of moderate or severe hepatic impairment in the last 365 days?					
4.	4. Has the patient tried 90 days of atorvastatin?				🗌 No	
5.	5. Has the patient tried 90 days of rosuvastatin? Yes					
6.	6. Has the patient tried 90 days of treatment with ezetimibe concurrently with atorvastatin or rosuvastatin,					
	immediately prior to PCSK9 inhibitor PA request?					
7.	7. Is the low density lipoprotein-cholesterol (LDL-C) level >70mg/dl despite treatment with 90 days of					
	atorvastatin treatment, 90	days of rosuvastatin, and most recently, 90 days of ez	etimibe t	reatment? 🗌 Yes	🗌 No	
Section 2. Laboratory Information:						

LDL-C prior to initiation of PCSK9 treatment:		Date level obtained:
mg/dL		(for first time requests, level must be from previous 60 days)
Current LDL-C:	mg/dL*	Date level obtained: (level must be from previous 60 days)

*Required for renewal requests only. Must have at least a 50% reduction in LDL-C compared to LDL-C level prior to PCSK9 treatment initiation for patients with HeFH and at least a 30% reduction in LDL-C for patients with HoFH for renewal approval.

By signing below, I, the prescriber, certify that the information provided above is verifiable and accurate to the best of my knowledge.

Prescriber Signature:	Date:				
Prior Authorization of Benefits is not the practice of medicine or the substitute for the independent medical judgment of a treating physician. Only a treating physician can determine what medications are appropriate for a patient. Please refer to the applicable plan for the detailed information regarding benefits, conditions, limitations, and exclusions. The submitting provider certifies that the information provided is true, accurate, and complete and the requested services are medically indicated and necessary to the health of the patient. Note: Payment is subject to member eligibility. Authorization does not guarantee payment.					
Please fax or mail this form to: Prime Therapeutics LLC, Clinical Review Department	CONFIDENTIALITY NOTICE: This communication is intended only for the use of the individual entity to which it is addressed and may contain information that is privileged or confidential. If the reader of this message is				
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Eagan, Minnesota 55121	not the intended recipient, you are hereby notified that any dissemination, distribution or copying of this communication is strictly prohibited. If you have received this communication in error, please notify the sender immediately by				
TOLL FREE	telephone at 866.202.3474 and return the original message to Prime				
Fax: 877.243.6930 Phone: 855.457.0407	Therapeutics via U.S. Mail. Thank you for your cooperation.				