LIPOTROPICS, OTHER PRIOR AUTHORIZATION FORM







(form effective 1/6/2025)

Fax to PerformRx SM at **1-855-851-4058**, or to speak to a representative call **1-888-674-8720**.

DDIOD ALITHODIZ	ATION DEGL	EST INFORMATION					
		EST INFORMATION					
□ New request □ Renewal request □ Total # of pages:			Contact's phone number:				
Name of office contact:	cility contact/phone:						
PATIENT INFORMA	ATION						
Patient name:			Patient ID #:		DOB:		
Street address:							
Apt #: City/state/zip:				Phone:			
PRESCRIBER INFO	RMATION						
Prescriber name:							
Specialty:			NPI:			State license #:	
Street address:							
Suite #:	City/state/zip:						
Phone:			Fax:				
CLINICAL INFORM	ATION						
Medication requested:							
Preferred:				Non-Preferred:			
☐ Cholestyramine Powder		☐ Fenofibrate Nanocrystalized 48 mg Tablet (generic Tricor) ☐ Fenofibrate Nanocrystalized 145 mg Tablet (generic Tricor) ☐ Fenofibric Acid (Choline) DR 45 mg Capsule (generic Trilipix) ☐ Fenofibric Acid (Choline) DR 135 mg Capsule]]]	☐ Antara Capsule		☐ Fenofibric Acid 105 mg Tablet	
☐ Cholestyramine Powder Packet				☐ Colesevelam Powder Packet		(generic Fibricor)	
☐ Cholestyramine Light Powder				☐ Colesevelam Tablet		□ Fenoglide Tablet	
☐ Cholestyramine Light Powder Packet				☐ Colestid Granule		☐ Fibricor Tablet	
 □ Colestipol Tablet □ Ezetimibe Tablet □ Fenofibrate 54 mg Tablet (generic Lofibra Tablet) □ Fenofibrate 160 mg Tablet 				☐ Colestid Tablet		 ☐ Icosapent Ethyl Capsule (generic Vascepa) 	
				☐ Colestipol Granule		☐ Juxtapid Capsule	
				☐ Colestipol Granule Packet		☐ Leqvio Syringe	
				□ Evkeeza Vial		□ Lipofen Capsule	
(generic Lofibra Tablet)	iot	(generic Trilipix)		☐ Fenofibrate 50 mg Capsule (generic Lipofen)		□ Lopid Tablet	
☐ Fenofibrate Micronized		☐ Gemfibrozil Tablet		☐ Fenofibrate 150 mg Capsule (generic Lipofen) ☐ Fenofibrate 40 mg Tablet		□ Lovaza Capsule	
(generic Antara)		☐ Nexletol Tablet				□ Niacin ER Tablet (generic Niaspan)	
☐ Fenofibrate Micronized 1 (generic Antara)	130 mg Capsule	☐ Nexlizet Tablet				☐ Questran Powder	
☐ Fenofibrate Micronized (generic Lofibra Capsul	le)	 ☐ Omega-3 Ethyl Esters Capsule (generic Lovaza) ☐ Praluent Pen 		(generic Fenoglide) □ Fenofibrate 120 mg Tablet		☐ Questran Powder Packet	
				(generic Fenoglide)		☐ Questran Light Powder	
☐ Fenofibrate Micronized (generic Lofibra Capsul		☐ Prevalite Powder	☐ Fenofibrate (Micronized) 90 mg Capsule (generic Antara)			☐ Tricor Tablet	
☐ Fenofibrate Micronized 2	200 mg Capsule	☐ Prevalite Powder Packet		90 mg Capsule (generic Antara)		☐ Trilipix DR Capsule	
(generic Lofibra Capsul		☐ Repatha Sureclick		☐ Fenofibric Acid 35 mg Tablet		□ Welchol Powder Packet	
		· · · · · · · · · · · · · · · · · · ·		(generic Fibricor)		□ Welchol Tablet	
D (□ Zetia Tablet	
Dosage form:						Strength:	
Dose/directions: Quantity:						Refills:	
Diagnosis:					[Ox code <i>(required)</i> :	



N	NITIAL REQUESTS
	Complete all sections that apply to the beneficiary and this request. Check all that apply and <u>submit documentation</u> for each item.
١.	For treatment of ANY LIPID DISORDER: Has results of a lipid profile within the past 3 months (submit copy)
2.	For a PCSK9 INHIBITOR (eg, Leqvio, Praluent, Repatha), NEXLETOL (bempedoic acid), or NEXLIZET (bempedoic acid/ezetimibe):
	 □ One of the following related to history of statin use: □ Failed to achieve goal LDL-C or percentage reduction of LDL-C with maximally tolerated dose of ONE high-intensity statin (eg, atorvastatin, rosuvastatin) for at least THREE consecutive months □ List medications tried:
	 □ Is unable to tolerate high-intensity statins AND: □ Has a temporally related intolerance to high-intensity statins □ Tried and failed or has an intolerance to the lowest FDA-approved daily dose or alternate-day dosing of any statin for at least THREE months List medications tried:
	 ☐ Modifiable comorbid conditions that may enhance statin intolerance were ruled out and/or addressed by the prescriber (eg, drug interactions, hypothyroidism, vitamin D deficiency, etc.) ☐ Has a contraindication to statins Please explain:
	□ One of the following related to history of ezetimibe use: □ Failed to achieve goal LDL-C or percentage reduction of LDL-C with ezetimibe in combination with maximally tolerated dose of the highest-tolerated intensity statin (eg, atorvastatin, rosuvastatin) for at least THREE consecutive months □ Has a contraindication or an intolerance to ezetimibe Please explain:
	For a PCSK9 inhibitor, has an LDL-C that is >25% above goal LDL-C while adherent to treatment with the maximally tolerated dose of the highest-tolerated intensity statin for at least THREE consecutive months List medications tried:
	□ One of the following: □ For a diagnosis of homozygous familial hypercholesterolemia, is prescribed the requested medication in addition to other standard lipid-lowering therapies □ For all other diagnoses, is prescribed the requested medication in addition to the maximally tolerated dose of the highest-tolerated intensity statin (if clinically appropriate) □ For a non-preferred PCSK9 inhibitor:
	☐ Tried and failed a preferred PCSK9 inhibitor or has a contraindication or an intolerance to the preferred PCSK9 inhibitors approved or medically accepted for the treatment of the beneficiary's diagnosis (Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.) List medications tried:
	□ For Nexletol (bempedoic acid) or Nexlizet (bempedoic acid/ezetimibe): □ If currently taking simvastatin or pravastatin, will not be using Nexletol/Nexlizet concomitantly with simvastatin at a dose of >20 mg daily or pravastatin at a dose of >40 mg daily
3.	For EVKEEZA (evinacumab) or JUXTAPID (lomitapide): Is prescribed the requested medication by or in consultation with a cardiologist, endocrinologist, or other provider specializing in lipid disorders Has a diagnosis of homozygous familial hypercholesterolemia in accordance with current consensus guidelines One of the following: Tried and failed or has a contraindication or an intolerance to PCSK9 inhibitors
	Please explain: □ Is homozygous for LDL receptor (LDLR)-negative mutations (ie, has LDLR-negative mutations in both alleles) associated with LDLR activity below 2% □ Is prescribed the requested medication in addition to other standard lipid-lowering therapies
l.	For VASECPA (icosapent ethyl):
	☐ Has a contraindication to statins Please explain:
5.	For ALL OTHER NON-PREFERRED Lipotropics, Other: Tried and failed or has a contraindication or an intolerance to the preferred Lipotropics, Other approved or medically accepted for the beneficiary's diagnosis (Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.) List medications tried:

RENEWAL REQUESTS

1. For ALL diagnoses:

☐ Experienced a positive clinical response demonstrated by lab test results, if appropriate for the diagnosis, since starting the requested medication (e.g., decreased LDL-C, decreased triglycerides, etc.) (submit copy of results)

2. For a PCSK9 INHIBITOR (eg, Leqvio, Praluent, Repatha):

- ☐ For a diagnosis of homozygous familial hypercholesterolemia, is using the requested PCKS9 inhibitor in addition to other standard lipid-lowering treatments
- ☐ For all other diagnoses, is using the requested PCSK9 inhibitor in addition to the maximally tolerated dose of the highest-tolerated intensity statin (if clinically appropriate)

3. For NEXLETOL (bempedoic acid) or NEXLIZET (bempedoic acid/ezetimibe):

- ☐ Is using the requested medication in addition to the maximally tolerated dose of the highest-tolerated intensity statin (if clinically appropriate)
- ☐ If currently taking simvastatin or pravastatin, will not be using Nexletol/Nexlizet concomitantly with simvastatin at a dose of >20 mg daily or pravastatin at a dose of >40 mg daily

4. For EVKEEZA (evinacumab) or JUXTAPID (lomitapide):

- ☐ Is prescribed the requested medication by or in consultation with a cardiologist, endocrinologist, or other provider specializing in lipid disorders
- ☐ Is using the requested medication in addition to other standard lipid-lowering treatments

5. For ALL OTHER NON-PREFERRED Lipotropics, Other:

☐ Tried and failed or has a contraindication or an intolerance to the preferred Lipotropics, Other approved or medically accepted for the beneficiary's diagnosis (Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.)

List medications tried:

Prescriber signature: Date:

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