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Section:	Prescription Drugs	**Effective Date:**	April 1, 2026
Subsection:	Cardiovascular Agents	**Original Policy Date:**	December 19, 2025
Subject:	Redemplo	**Page:**	1 of 4

Last Review Date: March 6, 2026

Redemplo

Description

Redemplo (plozasiran)

Background
Redemplo (plozasiran) is a small interfering ribonucleic acid (siRNA) conjugated with GalNAc that degrades the apolipoprotein C-III (apoC-III) mRNA through the RNA interference mechanism resulting in reduced levels of hepatic and serum apoC-III protein. Reduction of apoC-III protein leads to increased clearance of serum triglycerides (1).

Regulatory status
FDA-approved indication: Redemplo is an apolipoprotein C-III (apoC-III)-directed small interfering ribonucleic acid (siRNA) indicated as an adjunct to diet to reduce triglycerides in adults with familial chylomicronemia syndrome (FCS) (1).

The safety and effectiveness of Redemplo in pediatric patients less than 18 years of age have not been established (1).

Related policies
Tryngolza

Policy

This policy statement applies to clinical review performed for pre-service (Prior Approval, Precertification, Advanced Benefit Determination, etc.) and/or post-service claims.

Redemplo may be considered **medically necessary** if the conditions indicated below are met.

Redemplo may be considered **investigational** for all other indications.

Prior-Approval Requirements

Age 18 years of age or older

Diagnosis

Patient must have the following:

Familial chylomicronemia syndrome (FCS)

AND ALL of the following:

1. Patient has **ONE** of the following:
    a. Diagnosis confirmed by genetic testing documenting biallelic pathogenic variants in FCS-causing genes (e.g., LPL, GPIHBP1, APOA5, APO2, LMF1, GPD1, CREB3L3)
    b. North American familial syndrome (NAFCS) score ≥45 **OR** Moulin score ≥10
    (e.g., *https://tryngolzahcp.com/diagnosing-fcs/fcs-diagnostic-scoring?tab=nafcs, https://tryngolzahcp.com/diagnosing-fcs/fcs-diagnostic-scoring?tab=moulin*)
2. Fasting triglyceride (TG) levels ≥880 mg/dL
3. Used in combination with a low-fat diet (≤20g of fat per day)
4. **NO** dual therapy with Tryngolza (olezarsen)

Prior – Approval *Renewal* Requirements

Age 18 years of age or older

Diagnosis

Patient must have the following:

Familial chylomicronemia syndrome (FCS)

AND ALL of the following:

1. Reduction in fasting triglycerides from baseline

2. Used in combination with a low-fat diet (≤20g of fat per day)
3. **NO** dual therapy with Tryngolza (olezarsen)

Policy Guidelines

Pre - PA Allowance
None

Prior - Approval Limits

Quantity 1 prefilled syringe per 84 days

Duration 12 months

Prior – Approval *Renewal* Limits
Same as above

Rationale

Summary
Redemplo is an apoC-III-directed siRNA indicated to reduce triglycerides in adults with familial chylomicronemia syndrome (FCS). The safety and effectiveness of Redemplo in pediatric patients less than 18 years of age have not been established (1).

Prior authorization is required to ensure the safe, clinically appropriate, and cost-effective use of Redemplo while maintaining optimal therapeutic outcomes.

References
1. Redemplo [package insert]. Pasadena, CA: Arrowhead Pharmaceuticals, Inc.; November 2025.

Policy History

Date	Action
December 2025	Addition to PA
March 2026	Annual review. Per SME, added option of NAFCS or Moulin score instead of genetic confirmation

Keywords

This policy was approved by the FEP® Pharmacy and Medical Policy Committee on March 6, 2026 and is effective on April 1, 2026.