

Promise Health Plan

inclisiran (Leqvio)

Medical Benefit Drug Policy

Place of Service

Home Infusion Administration Infusion Center Administration

Office Administration

Outpatient Facility Administration

Drug Details

USP Category: CARDIOVASCULAR AGENTS

Mechanism of Action: Small interfering RNA (siRNA) directed to PCSK9 mRNA

HCPCS:

J1306:Injection, inclisiran, 1 mg

How Supplied:

284 mg/1.5 mL (189 mg/mL) single-dose prefilled syringe

Condition(s) listed in policy (see coverage criteria for details)

 Clinical Atherosclerotic Cardiovascular Disease (ASCVD), Heterozygous Familial Hypercholesterolemia (HeFH), Prevention of CVD Events, or Hypercholesterolemia

Any condition not listed in this policy requires a review to confirm it is medically necessary. For conditions that have not been approved for intended use by the Food and Drug Administration (i.e., off-label use), the criteria outlined in the California Code of Regulations (CCR), Title 22, Section 51303 and 51313 must be met.

Special Instructions and Pertinent Information

Provider must submit documentation (such as office chart notes, lab results or other clinical information) to ensure the member has met all medical necessity requirements.

The member's specific benefit may impact drug coverage. Other utilization management processes, and/or legal restrictions may take precedence over the application of this clinical criteria.

Coverage Criteria

The following condition(s) require Prior Authorization/Preservice.

<u>Clinical Atherosclerotic Cardiovascular Disease (ASCVD), Heterozygous Familial</u> <u>Hypercholesterolemia (HeFH), Prevention of CVD Events, or Hypercholesterolemia</u> Meets medical necessity if all the following are met:

- 7. Current LDL cholesterol (LDL-C) is at least 70 mg/dl (or at least 55 mg/dl) if extreme risk for heart disease), and one of the following (a, b, or c):
 - a. Patient has had at least 8 weeks of treatment with a high-intensity statin
 - b. Patient has an FDA-approved package insert (PI) supported contraindication to treatment with all statins

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- c. Patient has statin intolerance as evidenced by experiencing one of the following (i, ii, or iii):
 - i. Statin-related rhabdomyolysis
 - ii. Statin-related skeletal muscle symptoms (e.g., myopathy, myalgia)
 - iii. Statin-related elevated hepatic transaminase
- 2. Inadequate response or intolerable side effect to Praluent (alirocumab) or Repatha (evolocumab), or contraindication to both Praluent and Repatha
- 3. Dose does not exceed FDA approved maximum

Covered Doses:

Up to 284 mg given as a single subcutaneous injection initially, again at 3 months, and then every 6 months

Coverage Period:

Yearly, based on continued response to therapy

ICD-10:

E78.00, E78.2, E78.5

References

Effective: 12/01/2025

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- 2. DrugDex. Available by subscription at http://www.micromedexsolutions.com/home/dispatch
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- 4. Handelsman Y, Jellinger PS, Guerin CK, et al. Consensus Statement by the American Association of Clinical Endocrinologists and American College of Endocrinology on the Management of Dyslipidemia and Prevention of Cardiovascular Disease Algorithm 2020 Executive Summary. *Endocr Pract.* 2020;26(10):1196-1224.
- 5. Jellinger PS, Handelsman Y, Rosenblit PD, et al. American Association of Clinical Endocrinologists and American College of Endocrinology Guidelines for Management of Dyslipidemia and Prevention of Cardiovascular Disease. *Endocr Pract.* 2017;23(Suppl 2):1-87.
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- 7. Writing Committee; Lloyd-Jones DM, Morris PB, Ballantyne CM, et al. 2022 ACC Expert Consensus Decision Pathway on the Role of Nonstatin Therapies for LDL-Cholesterol Lowering in the Management of Atherosclerotic Cardiovascular Disease Risk: A Report of the American College of Cardiology Solution Set Oversight Committee. J Am Coll

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- 8. CMS Benefit Policy Manual. Chapter 15; § 50 Drugs and Biologicals Medicare Coverage Database. Available at https://www.cms.gov/Medicare-Coverage-Database/search.aspx
 Social Security Act (Title XVIII) Standard References, Sections: 1862(a)(1)(A) Medically Reasonable & Necessary; 1862(a)(1)(D) Investigational or Experimental; 1833(e) Incomplete Claim; 1861(t) (1) Drugs and Biologicals

Review History

Effective: 12/01/2025

Date of Last Annual Review: 4Q2025 Changes from previous policy version:

- Clarify coverage to include hypercholesterolemia. Rationale: In July 2025, the FDA expanded approval to use as first-line monotherapy for LDL-C reduction in adults with hypercholesterolemia
- Clarify coverage for statin related intolerance. Rationale: American College of Cardiology NLA 2022 Definition of Statin Intolerance

Blue Shield of California Medication Policy to Determine Medical Necessity Reviewed by P&T Committee

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