

RNA-Targeted Therapies (Amvuttra® and Onpattro®) (for Louisiana Only)

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[Instructions for Use](#)

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Application

This Medical Benefit Drug Policy only applies to the state of Louisiana.

Coverage Rationale

Amvuttra® (vutrisiran) and Onpattro® (patisiran) are proven and medically necessary for the treatment of the polyneuropathy of hereditary transthyretin-mediated (hATTR) amyloidosis in patients who meet all of the following criteria:

- For **initial therapy**, all of the following:
 - **Both** of the following:
 - Diagnosis of hATTR amyloidosis with polyneuropathy
 - Documentation that the patient has a pathogenic TTR mutation (e.g., V30M)
 - and**
 - Documentation of **one** of the following:
 - Patient has a baseline polyneuropathy disability (PND) score ≤ IIIb
 - Patient has a baseline familial amyloid polyneuropathy (FAP) Stage 1 or 2
 - Patient has a baseline neuropathy impairment score (NIS) ≥ 5 and ≤ 130
 - Patient has a baseline Karnofsky performance status (KPS) score ≥ 60%
 - and**
 - Presence of clinical signs and symptoms of the disease (e.g., peripheral sensorimotor polyneuropathy, autonomic neuropathy, motor disability, etc.); **and**
 - Patient has not had a liver transplant; **and**
 - **One** of the following:
 - For **Amvuttra**, patient is not receiving in combination with **any** of the following:
 - RNA interference agents [e.g., Onpattro (patisiran), Wainua (eplontersen), Tegsedi (inotersen)]
 - Transthyretin stabilizers [e.g., Vyndaqel (tafamidis meglumine) or Vyndamax (tafamidis)]
 - For **Onpattro**, patient is not receiving in combination with **any** of the following:
 - RNA interference agents [e.g., Amvuttra (vutrisiran), Wainua (eplontersen), Tegsedi (inotersen)]
 - Transthyretin stabilizers [e.g., Vyndaqel (tafamidis meglumine) or Vyndamax (tafamidis)]
 - and**
 - Prescribed by or in consultation with a neurologist; **and**
 - Dosing is in accordance with the US Food and Drug Administration prescribing information; **and**

- Initial authorization is for no more than 12 months
- For **continuation of therapy**, all of the following:
 - Patient has previously received treatment with Amvuttra or Onpattro
 - and**
 - Documentation of **one** of the following:
 - Patient continues to have a PND score \leq IIIb
 - Patient continues to have a FAP stage 1 or 2
 - Patient continues to have a NIS score \geq 5 and \leq 130
 - Patient continues to have a KPS score \geq 60%
 - and**
 - Documentation that the patient has experienced a positive clinical response to Amvuttra or Onpattro (e.g., improved neurologic impairment, motor function, quality of life, slowing of disease progression, etc.); **and**
 - **One** of the following:
 - For **Amvuttra**, patient is not receiving in combination with **any** of the following:
 - RNA interference agents [e.g., Onpattro (patisiran), Wainua (eplontersen), Tegsedi (inotersen)]
 - Transthyretin stabilizers [e.g., Vyndaqel (tafamidis meglumine) or Vyndamax (tafamidis)]
 - For **Onpattro**, patient is not receiving in combination with **any** of the following:
 - RNA interference agents [e.g., Amvuttra (vutrisiran), Wainua (eplontersen), Tegsedi (inotersen)]
 - Transthyretin stabilizers [e.g., Vyndaqel (tafamidis meglumine) or Vyndamax (tafamidis)]
 - Prescribed by or in consultation with a neurologist; **and**
 - Dosing is in accordance with the US Food and Drug Administration prescribing information; **and**
 - Authorization is for no more than 12 months

Amvuttra® (vutrisiran) and Onpattro® (patisiran) are unproven and not medically necessary for the treatment of:

- Sensorimotor or autonomic neuropathy not related to hATTR amyloidosis
- Primary or leptomeningeal amyloidosis
- Transthyretin (ATTR)-mediated amyloidosis with cardiomyopathy (ATTR-CM)

Applicable Codes

The following list(s) of procedure and/or diagnosis codes is provided for reference purposes only and may not be all inclusive. Listing of a code in this policy does not imply that the service described by the code is a covered or non-covered health service. Benefit coverage for health services is determined by federal, state, or contractual requirements and applicable laws that may require coverage for a specific service. The inclusion of a code does not imply any right to reimbursement or guarantee claim payment. Other Policies and Guidelines may apply.

HCPCS Code	Description
J0222	Injection, patisiran, 0.1 mg
J0225	Injection, vutrisiran, 1 mg

Diagnosis Code	Description
E85.1	Neuropathic hereditary amyloidosis

Background

Hereditary ATTR (hATTR) amyloidosis, formerly known as familial amyloid polyneuropathy, is a progressive, disabling, and life-threatening polyneuropathy affecting the peripheral and autonomic nervous system. This disease is an autosomal transmission disorder which is usually due to a point mutation of the transthyretin (TTR) gene. The disease is caused by misfolded transthyretin (TTR) protein that accumulates as amyloid fibrils in multiple organs, including the nerves, heart, and gastrointestinal tract.

Amvuttra (vutrisiran) and Onpattro (patisiran) are double-stranded small interfering RNAs (siRNAs) that target a sequence of mRNA conserved across wild-type and all TTR variants and can thereby degrade and reduce serum levels and protein deposits in tissues of both wild-type and mutated protein. It is formulated as lipid nanoparticles which direct it to the liver, the primary source of circulating TTR. Patisiran therapy is associated with observed lowering of TTR levels in both wild-type and mutant (V30M) forms of TTR.

A genetic testing service is available in the United States and Canada and a genetic counseling service is available in the

Clinical Evidence

A randomized, double-blind, placebo-controlled, phase III, global study (APOLLO) evaluated the efficacy and safety of patisiran in patients with hATTR amyloidosis with polyneuropathy. Adult patients 18 to 85 years of age were eligible for the study if the investigatory estimated survival to be ≥ 2 years, Neuropathy Impairment Score (NIS) of 5 to 130, and polyneuropathy disability score \leq IIIb. Patients were randomized 2:1 (n = 148:77) to receive either intravenous (IV) patisiran 0.3 mg/kg or placebo every 3 weeks. The primary endpoint was to determine the efficacy of patisiran at 18 months based on the difference in the change in modified NIS + 7 (a composite measure of motor strength, sensation, reflexes, nerve conduction, and autonomic function) between the patisiran and placebo groups. Secondary endpoints evaluated the effect of patisiran on Norfolk-Diabetic Neuropathy quality of life questionnaire score, nutritional status (as evaluated by modified body mass index), motor function (as measured by NIS-weakness and timed 10-m walk test), and autonomic symptoms (as measured by the Composite Autonomic Symptom Score-31 questionnaire). Exploratory objectives include assessment of cardiac function and pathologic evaluation to assess nerve fiber innervation and amyloid burden. Safety of patisiran was also assessed throughout the study. Overall patisiran reduced the mean max serum TTR reduction by 87.8% from baseline in the patisiran treated group over 18 months. The LS mean change in the mNIS + 7 from baseline at 18 months was -33.99 (p = 9.26×10^{-24}); (patisiran -6.03; placebo + 27.96). The LS mean change in the Norfolk QOL-DN from baseline at 18 months was -21.1 (p = 1.10×10^{-10}); (patisiran -6.7; placebo + 14.4). All secondary endpoints (e.g., NIS-W, R-ODS, COMPASS-31, etc.) also achieved statistical significance at 18 months. The investigators also concluded that patisiran therapy was relatively safe and well tolerated with no increases in the frequency of events for patisiran compared to placebo group by system organ class. Overall, 13 deaths occurred in the APOLLO study, however, none of these were considered related to the study drugs and were consistent with natural history. The majority of infusion-related reactions were mild in severity, with no severe or life-threatening, or serious reactions. These reactions decreased over time and led to treatment discontinuation in only 1 patient. The investigators concluded that patisiran treatment resulted in significant improvement in polyneuropathy relative to placebo while significantly reducing disease symptoms and disability, improvement in quality of life, nutritional status, strength, and ambulation seen with patisiran relative to placebo.^{1,8}

In a subpopulation analysis of the APOLLO trial, investigators evaluated the treatment association of patisiran with regional left ventricular (LV) myocardial strain in cardiac manifestation in hATTR.^{11,12} The prespecified cardiac subpopulation [126 of 225 (56%)] comprised of patients with a baseline LV wall thickness of 13 mm or more and no history of hypertension or aortic valve disease. Of the 126 patients included in the prespecified cardiac subpopulation, 36 patients (28.6%) received placebo and 90 patients (71.4%) received patisiran. At baseline, LV global longitudinal strain (GLS) was impaired and regional longitudinal strains were lowest in the basal segments with apical sparing. There were no differences in regional longitudinal strains between the treatment groups at baseline. patisiran improved the absolute GLS [least-squares mean (SE) difference, 1.4% (0.6%); 95% CI, 0.3%-2.5%; p = .02] compared with placebo at 18 months, with the greatest differential increase observed in the basal region [overall least-squares mean (SE) difference, 2.1% (0.8%); 95% CI, 0.6%-3.6%; p = .006] and no significant differences in the mid and apical regions among groups. patisiran reduced mean left ventricular wall thickness (least-squares mean difference \pm SEM: -0.9 ± 0.4 mm, p = 0.017), interventricular septal wall thickness, posterior wall thickness, and relative wall thickness at month 18 compared with placebo. patisiran also led to increased end-diastolic volume (8.3 ± 3.9 mL, p = 0.036), decreased global longitudinal strain ($-1.4 \pm 0.6\%$, p = 0.015), and increased cardiac output (0.38 ± 0.19 L/min, p = 0.044) compared with placebo at month 18. patisiran lowered N-terminal prohormone of brain natriuretic peptide at 9 and 18 months (at 18 months, ratio of fold-change patisiran/placebo 0.45, p < 0.001). A consistent effect on N-terminal prohormone of brain natriuretic peptide at 18 months was observed in the overall APOLLO patient population (n = 225). Median follow-up duration was 18.7 months. The exposure-adjusted rates of cardiac hospitalizations and all-cause death were 18.7 and 10.1 per 100 patient-years in the placebo and patisiran groups, respectively (Andersen-Gill hazard ratio, 0.54; 95% CI, 0.28-1.01). The authors concluded that patisiran prevented the deterioration of LV GLS and decreased mean LV wall thickness over 18 months, suggesting that patisiran may halt or reverse the progression of the cardiac manifestations of hATTR amyloidosis.

The safety and efficacy of vutrisiran was established in a phase 3 randomized, open-label study (NCT03759379) in adult patients with polyneuropathy caused by hATTR amyloidosis. Patients eligible for the study met the following criteria: aged 18-25 years, diagnosis of ATTRv amyloidosis with a documented TTR variant and baseline NIS 5-130, polyneuropathy disability (PND) score \leq IIIb, a KPS ≥ 60 , and adequate liver and renal function. Patients were randomized 3:1 to receive 25 mg of vutrisiran subcutaneously once every 3 months (n = 122), or 0.3mg/kg patisiran intravenously every 3 weeks (n =) as a reference group. Efficacy assessments were based on a comparison of the vutrisiran with an external placebo group in another study (NCT01960348) composed of a comparable population of adult patients with polyneuropathy caused by hATTR amyloidosis. The primary endpoint was the change from baseline to month 9 in modified Neuropathy Impairment Score + 7 (mNIS + 7). The mNIS + 7 has a total score range from 0 to 304 points, with higher scores

representing a greater severity of disease. The least squares mean change from baseline for the mNIS + 7 score was -2.2 for vutrisiran vs. + 14.8 for placebo (difference of -17.0, 95% CI: -21.8, -12.2; $p < 0.001$). The clinical meaningfulness of effects on the mNIS + 7 was assessed by the change from baseline to Month 9 in Norfolk Quality of Life-Diabetic Neuropathy (QoL-DN) total score. The Norfolk QoL-DN scale is a patient-reported assessment that evaluates the subjective experience of neuropathy in the following domains: physical functioning/large fiber neuropathy, activities of daily living, symptoms, small fiber neuropathy, and autonomic neuropathy. The Norfolk QoL-DN has a total score range from -4 to 136, with higher scores representing greater impairment. Additional endpoints were gait speed, as measured by the 10-meter walk test (10MWT), and modified body mass index (mBMI). The mean least squares mean change from baseline for the Norfolk QoL-DN total score was -3.3 for vutrisiran vs. + 12.9 for placebo (difference of -16.2, 95% CI: -21.7, -10.8; $p < 0.001$). The mean least squares mean change from baseline for the 10-meter walk test was 0 for vutrisiran vs. -0.13 for placebo (difference of 0.13, 95% CI: 0.07, 0.19; $p < 0.001$) and 10-meter walk test at Month 9 compared to placebo in the external study ($p < 0.001$). The mean least squares mean change from baseline for mBMI was 7.6 for vutrisiran vs. -60.2 for placebo (difference of 67.8, 95% CI: 43.0, 92.6; $p < 0.001$).

The most common adverse reactions (at least 5%) were arthralgia (11%), dyspnea (7%), and decreased vitamin A (7%). Patients were instructed to take the recommended daily allowance of vitamin A. Seventy-four percent of patients treated with vutrisiran had normal vitamin A levels at baseline, and 98% of those with a normal baseline developed low vitamin A levels. In some cases, the decreased vitamin A level was reported as an adverse reaction. Two serious adverse reactions of atrioventricular (AV) heart block (1.6%) occurred in patients treated with vutrisiran, including one case of complete AV block. Injection site reactions were reported in 5 (4%) patients treated with vutrisiran. Reported symptoms included bruising, erythema, pain, pruritus, and warmth. Injection site reactions were mild and transient.

Institute for Clinical and Economic Review (ICER)

On October 4, 2018, ICER released a clinical report entitled, “Inotersen and patisiran for Hereditary Transthyretin Amyloidosis: Effectiveness and Value”. ICER recommendations are as follows:¹³

- ICER judges the clinical evidence for patisiran to be “incremental” or “better”.
- On average, patients on patisiran demonstrated improvement in neuropathy symptoms, as measured by the mNIS + 7. Based on the current body of evidence, there is moderate certainty of a substantial net health benefit with high certainty of at least a small net health benefit compared to best supportive care.

U.S. Food and Drug Administration (FDA)

This section is to be used for informational purposes only. FDA approval alone is not a basis for coverage.

Amvuttra® (vutrisiran) is a transthyretin-directed small interfering RNA indicated for the treatment of the polyneuropathy of hereditary transthyretin-mediated amyloidosis in adults.

Onpattro® (patisiran) contains a transthyretin-directed small interfering RNA and is indicated for the treatment of the polyneuropathy of hereditary transthyretin-mediated amyloidosis in adults.

References

1. Onpattro [prescribing information]. Cambridge, MA: Alnylam Pharmaceuticals. January 2023.
2. Coutinho P, Martins da Silva A, Lopes Lima J, Resende Barbosa A. (1980) Forty years of experience with type I amyloid neuropathy. Review of 483 cases. In: Glenner G., Costa P., de Freitas A., editors. (eds), Amyloid and Amyloidosis. Amsterdam: Excerpta Medica, pp. 88–98.
3. Yamamoto S, Wilczek H, Nowak G, et al. Liver transplantation for familial amyloidotic polyneuropathy (FAP): a single-center experience over 16 years. *Am J Transplant*. 2007 Nov;7(11):2597-604.
4. Koike H, Misu K, Ikeda S, et al. Type I (transthyretin Met30) familial amyloid polyneuropathy in Japan: early- vs late-onset form. *Arch Neurol*. 2002 Nov;59(11):1771-6.
5. Koike H, Tanaka F, Hashimoto R, et al. Natural history of transthyretin Val30Met familial amyloid polyneuropathy: analysis of late-onset cases from non-endemic areas. *J Neurol Neurosurg Psychiatry*. 2012 Feb;83(2):152-8.
6. Adams D, Suhr OB, Dyck PJ, et al. Trial design and rationale for APOLLO, a Phase 3, placebo-controlled study of patisiran in patients with hereditary ATTR amyloidosis with polyneuropathy. *BMC Neurol*. 2017 Sep 11;17(1):181.
7. Adams D. Recent advances in the treatment of familial amyloid polyneuropathy. *Ther Adv Neurol Disord*. 2013 Mar; 6(2): 129–139.

8. Adams D, Gonzalez-Duarte A, O’Riordan WD, et al. patisiran, an RNAi Therapeutic, for Hereditary Transthyretin Amyloidosis. *N Engl J Med*. 2018 Jul 5;379(1):11-21.
9. Alnylam Pharmaceuticals. The Study of an Investigational Drug, patisiran (ALN-TTR02), for the Treatment of Transthyretin (TTR)-Mediated Amyloidosis in Patients Who Have Already Been Treated With ALN-TTR02 (patisiran). In: *ClinicalTrials.gov* [Internet]. Bethesda (MD): National Library of Medicine (US). 2000- [cited 2018 April 12]. Available at: <https://clinicaltrials.gov/show/NCT02510261>. NLM Identifier: NCT02510261.
10. Institute for Clinical and Economic Review: Draft Evidence Report - Inotersen and patisiran for Hereditary Transthyretin Amyloidosis: Effectiveness and Value. July 20, 2018.
11. Minamisawa M, Claggett B, Adams D, et al. Association of patisiran, an RNA Interference Therapeutic, With Regional Left Ventricular Myocardial Strain in Hereditary Transthyretin Amyloidosis: The APOLLO Study. *JAMA Cardiol*. 2019 Mar 16.
12. Solomon SD, Adams D, Kristen A, et al. Effects of patisiran, an RNA Interference Therapeutic, on Cardiac Parameters in Patients With Hereditary Transthyretin-Mediated Amyloidosis. *Circulation*. 2019 Jan 22;139(4):431-443.
13. Institute for Clinical and Economic Review: Inotersen and patisiran for Hereditary Transthyretin Amyloidosis: Effectiveness and Value. October 4, 2018.
14. Luigetti M, Romano A, Di Paolantonio A, Bisogni G, Sabatelli M. Diagnosis and Treatment of Hereditary Transthyretin Amyloidosis (hATTR) Polyneuropathy: Current Perspectives on Improving Patient Care. *Ther Clin Risk Manag*. 2020;16:109-123. Published 2020 Feb 21. doi:10.2147/TCRM.S219979.
15. Amvuttra [prescribing information]. Cambridge, MA: Alnylam Pharmaceuticals. February 2023.

Policy History/Revision Information

Date	Summary of Changes
11/01/2024	<p>Coverage Rationale</p> <ul style="list-style-type: none"> • Revised coverage criteria: <ul style="list-style-type: none"> ○ Updated list of examples of RNA interference agents the patient must not receive in combination with Amvuttra or Onpattro; added Wainua (eplontersen) <p>Onpattro (patisiran)</p> <ul style="list-style-type: none"> ○ Added criterion requiring: <ul style="list-style-type: none"> Initial Therapy <ul style="list-style-type: none"> ▪ The patient has a baseline Karnofsky performance status (KPS) score \geq 60% Continuation of Therapy <ul style="list-style-type: none"> ▪ The patient continues to have a KPS score \geq 60% • Added language to indicate: <ul style="list-style-type: none"> ○ Amvuttra (vutrisiran) is unproven and not medically necessary for the treatment of: <ul style="list-style-type: none"> ▪ Sensorimotor or autonomic neuropathy not related to hATTR amyloidosis ▪ Primary or leptomeningeal amyloidosis ▪ Transthyretin (ATTR)-mediated amyloidosis with cardiomyopathy (ATTR-CM) ○ Onpattro (patisiran) is unproven and not medically necessary for the treatment of transthyretin (ATTR)-mediated amyloidosis with cardiomyopathy (ATTR-CM) <p>Supporting Information</p> <ul style="list-style-type: none"> • Archived previous policy version CSLA2023D0072L

Instructions for Use

This Medical Benefit Drug Policy provides assistance in interpreting UnitedHealthcare standard benefit plans. When deciding coverage, the federal, state or contractual requirements for benefit plan coverage must be referenced as the terms of the federal, state or contractual requirements for benefit plan coverage may differ from the standard benefit plan. In the event of a conflict, the federal, state or contractual requirements for benefit plan coverage govern. Before using this policy, please check the federal, state or contractual requirements for benefit plan coverage. UnitedHealthcare reserves the right to modify its Policies and Guidelines as necessary. This Medical Benefit Drug Policy is provided for informational purposes. It does not constitute medical advice.

UnitedHealthcare may also use tools developed by third parties, such as the InterQual® criteria, to assist us in administering health benefits. The UnitedHealthcare Medical Benefit Drug Policies are intended to be used in connection

with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.