

*You are invited to attend a
presentation on*

ATTR-CM Management: A Detailed Overview of Attruby

Date

Wednesday, July 29, 2026

Time

06:00 PM EST

Leo's Seafood

60 Ottawa Avenue Northwest
Grand Rapids, MI 49503-2646

Presented by

Michael G Dickinson, MD
Advanced Heart Failure and Transplant
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Hosted by

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Registration

<https://rsvp.bridgebio.cm-go.com/home/index/BR-34088>

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INDICATION

Attruby® (acoramidis) is indicated for the treatment of the cardiomyopathy of wild-type or variant transthyretin-mediated amyloidosis (ATTR-CM) in adults to reduce cardiovascular death and cardiovascular-related hospitalization.

SELECT SAFETY INFORMATION

Diarrhea (11.6% vs 7.6%) and upper abdominal pain (5.5% vs 1.4%) were reported in patients treated with Attruby versus placebo, respectively. The majority of these adverse reactions were mild and resolved without drug discontinuation. Increase in serum creatinine and decrease in eGFR may occur within 4 weeks of starting Attruby and then stabilize. The laboratory changes were reversible after treatment discontinuation.

*Please see additional Important Safety Information on the reverse side and accompanying full **Prescribing Information**.*

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INDICATION

Attruby® (acoramidis) is indicated for the treatment of the cardiomyopathy of wild-type or variant transthyretin-mediated amyloidosis (ATTR-CM) in adults to reduce cardiovascular death and cardiovascular-related hospitalization.

IMPORTANT SAFETY INFORMATION

Adverse Reactions

Diarrhea (11.6% vs 7.6%) and upper abdominal pain (5.5% vs 1.4%) were reported in patients treated with Attruby versus placebo, respectively. The majority of these adverse reactions were mild and resolved without drug discontinuation.

Discontinuation rates due to adverse events were similar between patients treated with Attruby versus placebo (9.3% and 8.5%, respectively).

Laboratory Tests

Mean increase in serum creatinine of 0.2 and 0.0 mg/dL and a mean decrease in eGFR of 8.2 and 0.7 mL/min/1.73 m² was observed in the adults with ATTR-CM treated with Attruby versus placebo, respectively, at Day 28 and then stabilized. These changes were reversible after treatment discontinuation.

Use in Specific Populations

Pregnancy & Lactation: There are no data on the use of Attruby in pregnant women. Animal data have not shown developmental risk associated with the use of Attruby in pregnancy.

There are no available data on the presence of Attruby in either human or animal milk or the effects of the drug on the breastfed infant or maternal milk production.