Omega-3 Fatty Acid Products

Policy # 00336
Original Effective Date: 01/09/2013
Current Effective Date: 01/17/2018

Applies to all products administered or underwritten by Blue Cross and Blue Shield of Louisiana and its subsidiary, HMO Louisiana, Inc. (collectively referred to as the “Company”), unless otherwise provided in the applicable contract. Medical technology is constantly evolving, and we reserve the right to review and update Medical Policy periodically.

When Services May Be Eligible for Coverage
Coverage for eligible medical treatments or procedures, drugs, devices or biological products may be provided only if:
- Benefits are available in the member’s contract/certificate, and
- Medical necessity criteria and guidelines are met.

Based on review of available data, the Company may consider brand or generic prescription omega-3 fatty acid products, including, but not limited to Lovaza® (omega-3-acid ethyl esters capsules), Vascepa® (icosapent ethyl capsules), Epanova® (omega-3-carboxylic acid capsules), or Omtryg® (omega-3-acid ethyl esters A capsules) to be eligible for coverage when the following patient selection criteria are met:

Patient Selection Criteria
Coverage eligibility will be considered for brand or generic prescription omega-3 fatty acid products when the following criteria are met:
- The patient has a fasting baseline (pretreatment) triglyceride (TG) level of ≥ 150 mg/dL; and
- The patient has tried, or is currently receiving, one of the following products: niacin (immediate-release or extended-release), a fibrate (e.g., gemfibrozil, fenofibrate, fenofibric acid), a statin (e.g., atorvastatin, simvastatin), or an over-the-counter (OTC) omega-3 fatty acid product (e.g., fish oil supplements).

(Note: This specific patient criterion is an additional Company requirement for coverage eligibility and will be denied as not medically necessary if not met)

When Services Are Considered Investigational
Coverage is not available for investigational medical treatments or procedures, drugs, devices or biological products.

Based on review of available data, the Company considers the use of brand or generic prescription omega-3 fatty acid products for a baseline (TG) level of < 150 mg/dL, or for usage not included in the above patient selection criteria to be investigational (with the exception of those denoted above as not medically necessary).

When Services Are Considered Not Medically Necessary
Based on review of available data, the Company considers the use of brand or generic prescription omega-3 fatty acid products when the patient has not tried or is not currently receiving, one of the following products: niacin (immediate-release or extended-release), a fibrate (e.g., gemfibrozil, fenofibrate, fenofibric

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Omega-3 acid, a statin (e.g., atorvastatin, simvastatin), or an OTC omega-3 fatty acid product (e.g., fish oil supplements) to be not medically necessary.**

**Background/Overview**

These drugs are combinations of omega-3 fatty acids and are indicated as an adjunct to diet to reduce TG levels in adult patients with severe (≥ 500 mg/dL) hypertriglyceridemia.

**Rationale/Source**

Several organizations including the National Cholesterol Education Program (NCEP), the American Heart Association (AHA), American Association of Clinical Endocrinologists' (AACE), and the Endocrine Society have released statements and/or guidelines which conclude that fibrates have the highest TG lowering effects out of the potential lipid lowering drugs. Other drugs with TG lowering effects include niacin, statins, and omega-3 fatty acid products. In fact, fibrates are considered the first-line treatment for hypertriglyceridemia, and more specifically severe hypertriglyceridemia. Patient selection criteria are based on information collected in a review of the available data.

**References**


**Policy History**

Original Effective Date:  01/09/2013
Current Effective Date:  01/17/2018
01/03/2013   Medical Policy Committee review
01/09/2013   Medical Policy Implementation Committee approval. New policy.
01/09/2014   Medical Policy Committee review
01/15/2014   Medical Policy Implementation Committee approval. Added a Note to the Patient Selection Criteria stating that denials will be not medically necessary if criteria are not met. Made a reference to this not medically necessary Note in the investigational section. Coverage eligibility unchanged.
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01/08/2015  Medical Policy Committee review
01/21/2015  Medical Policy Implementation Committee approval. Added Omtryg and Epanova to policy.
01/07/2016  Medical Policy Committee review
01/22/2016  Medical Policy Implementation Committee approval. Added the term “brand or generic” now that there are generic products available. No coverage change.
01/05/2017  Medical Policy Committee review
01/18/2017  Medical Policy Implementation Committee approval. No coverage change.
01/04/2018  Medical Policy Committee review
01/17/2018  Medical Policy Implementation Committee approval. No coverage change.
Next Scheduled Review Date: 01/2019

*Investigational – A medical treatment, procedure, drug, device, or biological product is Investigational if the effectiveness has not been clearly tested and it has not been incorporated into standard medical practice. Any determination we make that a medical treatment, procedure, drug, device, or biological product is Investigational will be based on a consideration of the following:

A. whether the medical treatment, procedure, drug, device, or biological product can be lawfully marketed without approval of the U.S. Food and Drug Administration (FDA) and whether such approval has been granted at the time the medical treatment, procedure, drug, device, or biological product is sought to be furnished; or

B. whether the medical treatment, procedure, drug, device, or biological product requires further studies or clinical trials to determine its maximum tolerated dose, toxicity, safety, effectiveness, or effectiveness as compared with the standard means of treatment or diagnosis, must improve health outcomes, according to the consensus of opinion among experts as shown by reliable evidence, including:
   1. Consultation with the Blue Cross and Blue Shield Association technology assessment program (TEC) or other nonaffiliated technology evaluation center(s);
   2. credible scientific evidence published in peer-reviewed medical literature generally recognized by the relevant medical community; or
   3. reference to federal regulations.

**Medically Necessary (or “Medical Necessity”) - Health care services, treatment, procedures, equipment, drugs, devices, items or supplies that a Provider, exercising prudent clinical judgment, would provide to a patient for the purpose of preventing, evaluating, diagnosing or treating an illness, injury, disease or its symptoms, and that are:

A. in accordance with nationally accepted standards of medical practice;

B. clinically appropriate, in terms of type, frequency, extent, level of care, site and duration, and considered effective for the patient's illness, injury or disease; and

C. not primarily for the personal comfort or convenience of the patient, physician or other health care provider, and not more costly than an alternative service or sequence of services at least as likely to produce equivalent therapeutic or diagnostic results as to the diagnosis or treatment of that patient's illness, injury or disease.

For these purposes, "nationally accepted standards of medical practice" means standards that are based on credible scientific evidence published in peer-reviewed medical literature generally recognized by the relevant medical community, Physician Specialty Society recommendations and the views of Physicians practicing in relevant clinical areas and any other relevant factors.

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