# SmartPA Criteria Proposal

**Drug/Drug Class:** Triglyceride Lowering Agents PDL Edit  
**First Implementation Date:** December 6, 2006  
**Proposed Date:** September 17, 2020  
**Prepared For:** MO HealthNet  
**Prepared By:** MO HealthNet/Conduent  
**Criteria Status:** ☒ Existing Criteria

## Executive Summary

**Purpose:** The MO HealthNet Pharmacy Program will implement a state-specific preferred drug list.

**Why Issue Selected:** The fibric acid derivatives (fibrates) are designed to lower triglycerides and to raise high-density lipoprotein (HDL). Fibrates have been shown to lower triglycerides by as much as 35-50% and to raise HDL by as much as 15-25%. Omega-3-Fatty Acids are present in fatty fish such as mackerel, lake trout, herring sardines, albacore tuna, and salmon and also in dietary supplements of fish oil concentrates, but large quantities of these products are required to reach therapeutic doses. Lovaza is a highly purified ethyl ester concentrate of eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA), two kinds of omega-3-fatty acids. The major effect of Lovaza is to lower TGs, although the mechanism of action is not entirely understood. These agents should be used in addition to a diet restricted in saturated fat and cholesterol when the individual response has been inadequate.

Total program savings for the PDL classes will be regularly reviewed.
**Program-Specific Information:**

<table>
<thead>
<tr>
<th>Preferred Agents</th>
<th>Non-Preferred Agents</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Fenofibrate 54, 67, 134, 160, 200mg (gen Lofibra®)</td>
<td>• Antara®</td>
</tr>
<tr>
<td>• Fenofibrate 48, 145mg (gen Tricor®)</td>
<td>• Fenofibrate (gen Antara®)</td>
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<tr>
<td>• Gemfibrozil</td>
<td>• Fenofibrate (gen Fenoglide®)</td>
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</tbody>
</table>

**Approval Criteria**
- Failure to achieve desired therapeutic outcomes with trial on 2 or more preferred agents
  - Documented trial period for preferred agents  **OR**
  - Documented ADE/ADR to preferred agents  **OR**
  - For Lovaza or Vascepa therapy: Triglyceride levels ≥ 500mg/dl allow first line therapy access

**Denial Criteria**
- Lack of adequate trial on required preferred agents
- Therapy will be denied if all approval criteria are not met

**Required Documentation**

<table>
<thead>
<tr>
<th>Laboratory Results:</th>
<th>Progress Notes:</th>
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<tbody>
<tr>
<td>X</td>
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**Type of Criteria:**  
- ☐ Increased risk of ADE  
- ☒ Preferred Drug List  
- ☐ Appropriate Indications  
- ☐ Clinical Edit  

**Data Sources:**  
- ☒ Only Administrative Databases  
- ☐ Databases + Prescriber-Supplied  

**Setting & Population**
- Drug class for review: Triglyceride Lowering Agents
- Age range: All appropriate MO HealthNet participants
Disposition of Edit

Denial: Exception Code "0160" (Preferred Drug List)
Rule Type: PDL

Default Approval Period

1 year

References

4. USPDI, Micromedex; 2020.
5. Facts and Comparisons eAnswers (online); 2020 Clinical Drug Information, LLC.