

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:	<input checked="" type="checkbox"/> Existing Criteria <input type="checkbox"/> Revision of Existing Criteria <input type="checkbox"/> New 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:	Preferred Agents	Non-Preferred Agents
	<ul style="list-style-type: none"> • Fenofibrate 54, 67, 134, 160, 200mg (gen Lofibra®) • Fenofibrate 48, 145mg (gen Tricor®) • Gemfibrozil 	<ul style="list-style-type: none"> • Antara® • Fenofibrate (gen Antara®) • Fenofibrate (gen Fenoglide®) • Fenofibrate (gen Lipofen®) • Fenofibrate (gen Triglide®) • Fenofibric Acid (gen Fibracor®) • Fenofibric Acid (gen Trilipix®) • Fenoglide® • Lipofen® • Lofibra® • Lopid® • Lovaza® • Omega-3 Acid Ethyl Esters • Tricor® • Triglide® • Triko • Trilipix® • Vascepa®

- 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

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 \geq 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

Laboratory Results:	<input checked="" type="checkbox"/>	Progress Notes:	<input type="checkbox"/>
MedWatch Form:	<input type="checkbox"/>	Other:	<input type="checkbox"/>

Disposition of Edit

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

Default Approval Period

1 year

References

1. Evidence-Based Medicine and Fiscal Analysis: "Triglyceride Lowering Agents – Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond, VA; July 2020.
2. Evidence-Based Medicine Analysis: "Triglyceride Lowering Agents", UMKC-DIC; June 2020.
3. Lippincott, Williams, Wilkins. PDR Electronic Library, Montvale NJ; 2020.
4. USPDI, Micromedex; 2020.
5. Facts and Comparisons eAnswers (online); 2020 Clinical Drug Information, LLC.

SmartPA PDL Proposal Form

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