



SmartPA Criteria Proposal

Drug/Drug Class:	Insulin, Rapid Acting PDL Edit		
First Implementation Date:	July 3, 2008		
Proposed Date:	July 18, 2023		
Prepared For:	MO HealthNet		
Prepared By:	MO HealthNet/Conduent		
Criteria Status:	☑ Existing Criteria☐ Revision of Existing Criteria☐ New Official		
	☐ New Criteria		

Executive Summary

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

Why Issue Selected:

Type 1 diabetes mellitus occurs when the body's immune system destroys the insulinsecreting beta cells of the pancreas. The management of type 1 diabetes has changed dramatically over the past 30 years. New insulin strategies have improved the ability to maintain near-normal glycemia. All rapid-acting insulins have demonstrated ability to lower hemoglobin A1c. An inhaled insulin product (Afrezza®) is now also available as part of this class but is indicated for adults only. Additional adverse effects of Afrezza include cough and throat pain and it is contraindicated with chronic lung diseases such as COPD or asthma. Factors such as onset, peak, and duration of action can influence the ability of an insulin regimen to help control glucose levels. Patient factors, including individual variations in insulin absorption, levels of exercise and types of meals consumed, also influence the effectiveness of insulin regimens.

Total program savings for the PDL classes will be regularly reviewed.

Program-Specific Information:

Preferred Agents	Non-Preferred Agents		
Humalog® Cartridge/Vial	Admelog® SoloStar® Pen/Vial		
 NovoLog® Cartridge/FlexPen®/Vial 	Afrezza® Cartridge		
	Apidra® SoloStar® Pen/Vial		
	Fiasp® FlexTouch®/PenFill®/Vial		
	Humalog KwikPen®, Tempo™ Pen		
	Humalog® Jr KwikPen®		
	 Insulin Aspart FlexPen®/PenFill®/Vial 		
	Insulin Lispro Jr KwikPen®		
•	 Insulin Lispro KwikPen®/Vial 		
	Lyumjev [®]		

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: Insulin, Rapid Acting
- 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 AND
- For insulin lispro 200 units/mL: documented compliance on prior rapid acting insulin therapy (90/120 days)

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Deniai Criteria
 Lack of adequate trial on required preferred agents Therapy will be denied if all approval criteria are not met
Required Documentation
Laboratory Results: Progress Notes: Other:
Disposition of Edit
Denial: Exception Code "0160" (Preferred Drug List) Rule Type: PDL
Default Approval Period

1 year

References

- Evidence-Based Medicine and Fiscal Analysis: "Therapeutic Class Review: ENDOCRINE AND METABOLIC AGENTS: Insulins, Rapid Acting", Gainwell Technologies; Last updated April 12, 2023.
- Evidence-Based Medicine Analysis: "Endocrine and Metabolic Agents: Rapid Acting", UMKC-DIC; February 2023.
- USPDI, Micromedex; 2023.
- Clinical Pharmacology [online]. Tampa (FL): Elsevier. 2023.