



SmartPA Criteria Proposal

Drug/Drug Class:	Beta Adrenergic Blockers and Beta Adrenergic Blockers/Diuretic Combinations PDL Edit	
First Implementation Date:	July 19, 2004	
Proposed Date:	September 16, 2021	
Prepared for:	MO HealthNet	
Prepared by:	MO HealthNet/Conduent	
Criteria Status:	⊠Existing Criteria □Revision of Existing Criteria □New Criteria	

Executive Summary

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

Why was this Issue Selected:

Beta-adrenergic blockers inhibit the chronotropic, inotropic and vasodilator responses to adrenaline by blocking β_1 and β_2 receptor sites throughout the body. Several characteristics of beta-blockers may be related to their clinical effectiveness. Beta blockers can be classified by cardioselectivity and intrinsic sympathomimetic activity (ISA). Cardioselective beta-blockers preferentially inhibit only β_1 receptors that are principally found in the myocardium. Non-cardioselective beta blockers inhibit both β_1 and β_2 receptor sites.

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

Program-specific information:

Preferred Agents	Non-Preferred Agents
Acebutolol	Betapace®
Atenolol	Betapace AF®
Atenolol/Chlorthalidone	Betaxolol
Bisoprolol	Bystolic®
Bisoprolol/HCTZ	Carvedilol ER
Carvedilol	Coreg®
Hemangeol®	Coreg CR®
Labetalol	Corgard®
Metoprolol Succinate	Inderal LA®
Metoprolol Tartrate	Inderal XL®
Metoprolol/HCTZ	InnoPran XL®
Nadolol	Kapspargo® Sprinkle Caps
Propranolol Soln/Tabs	Lopressor®
Propranolol/HCTZ	Lopressor HCT®
Sorine®	Nadolol/Bendroflumethiazide
Sotalol	Pindolol
Sotalol AF	Propranolol LA
	Sotylize®

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		 Tenoretic® Tenormin® Timolol Maleate Toprol XL® Ziac® 		
Type of Criteria:	☐ Increased risk of ADE☐ Appropriate Indications	☑ Preferred Drug List ☐ Clinical Edit		
Data Sources:	☐ Only Administrative Databases	☑ Databases + Prescriber-Supplied		
Setting & Popula	tion			
Combinations	s for review: Beta Adrenergic Blockers and appropriate MO HealthNet participants	Beta Adrenergic Blockers Diuretic		
Approval Criteria				
 Failure to achieve desired therapeutic outcomes with trial on 3 or more preferred agents Documented trial period for preferred agents OR Documented ADE/ADR to preferred agents For Coreg CR: Documented diagnosis of heart failure AND Adequate therapeutic trial on carvedilol twice daily for 30 days For Bystolic: Adequate therapeutic trial on one vasodilating alpha/beta-adrenergic blocking agent (labetalol or carvedilol) For Hemangeol: Participants aged 2 years and younger AND Documented diagnosis of infantile hemangioma AND Maximum treatment length of 6 months; clinical consultant review required to extended treatment For Sotylize and Kapspargo Sprinkle: Clinical Consultant Review for participants aged 10 years or older 				
Denial Criteria				
•	ate trial on required preferred agents denied if all approval criteria are not met			
Required Docum	entation			
Laboratory Result MedWatch Form:	Progress Notes: Other:			
Disposition of Ed	lit			
Denial: Exception Rule Type: PDL	Code "0160" (Preferred Drug List Edit)			

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Default Approval Period

1 year

References

- Evidence-Based Medicine and Fiscal Analysis: "Beta Adrenergic Blockers and Diuretic Combination Agents – Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond, VA; July 2021.
- 2. Evidence-Based Medicine Analysis: "Oral Beta Blockers (and any beta-blocker combinations)", UMKC-DIC; November 2003 August 2020.
- 3. Evidence-Based Medicine Analysis: "Beta Adrenergic Blockers and Diuretic Combinations", UMKC-DIC; June 2021.
- 4. USPDI, Micromedex; 2021.
- 5. Facts and Comparisons eAnswers (online); 2021 Clinical Drug Information, LLC.

