

Reference number(s) 6555-D

This document applies to the following:

Formulary	Applies
Standard Control (SF)	V
Standard Control - Choice (SCCF)	V
Preferred Drug Plan Design (PDPD)	
Advanced Control Specialty (ACSF)	V
Advanced Control Specialty - Choice (ACSCF)	V
Managed Medicaid Template (MMT)	
Marketplace (MF)	
Aetna Small Group Affordable Care Act (SG ACA) Aetna Health Exchange (AHE)	
Aetna Individual Lives (IVL)	
Value (VF)	V
New to Market (NTM)	

Formulary	Applies
Standard Formulary Chart (SFC)	
Basic Control Chart Preferred Drug Plan Design (BCC PDPD)	
Advanced Control Specialty Formulary Chart (ACSFC)	
Value Formulary Chart (VFC)	V
Medical Benefit	
Medical Benefit: Advanced Biosimilars First	
Combined Benefit Medical Specialty (CBMS)	
Medical Benefit: Managed Medicaid (MMMB)	
Medicare Part B	
Medicare Part B: Advanced Biosimilars First	

Exceptions Criteria Cushing's Syndrome

This document informs prescribers of preferred products and provides an exception process for the targeted product through prior authorization.

These criteria were developed to align with the following: Standard Control (SF), Standard Control – Choice (SCCF), Advanced Control Specialty Formulary (ACSF), Advanced Control Specialty – Choice Formulary (ACSCF), Value Formulary (VF), and Value Formulary Chart (VFC).

Plan Design Summary

This program applies to the Cushing's syndrome products specified in this document. Coverage for the targeted product is provided based on clinical circumstances that would exclude the use of the preferred product and may be based on previous use of a product. The coverage review process will ascertain situations where a clinical exception can be made. This program applies to all members requesting treatment with the targeted product.

Each referral is reviewed based on all utilization management (UM) programs implemented for the client.

Specialty Exceptions Cushing's Syndrome SF-SCCF-ACSF-ACSCF-VF-VFC 6555-D P2025a.docx © 2025 CVS Caremark. All rights reserved.

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Table. Cushing's Syndrome Products

Medications considered formulary or preferred on your plan may still require a clinical prior authorization review.

	Product(s)
Preferred	mifepristone (generic)
Target	Korlym (mifepristone)

Exception Criteria

This program applies to members requesting treatment for an indication that is FDA-approved for the preferred product.

Coverage for Korlym is provided when the member has had a documented intolerable adverse event to the preferred product generic mifepristone, and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information.

References

- 1. Korlym [package insert]. Redwood City, CA: Corcept Therapeutics Incorporated; September 2024.
- 2. Mifepristone [package insert]. Redwood City, CA: Corcept Therapeutics Incorporated; September 2024.