



## RX.PA.033.CCH SPECIALTY DRUG MANAGEMENT

The purpose of this policy is to define the prior authorization process for **specialty drugs processed under the medical benefit that do not have an existing drug specific policy.**

A specialty drug is any high-cost drug including injectables, infused products, oral agents, or inhaled medications, which require unique storage/ shipment and additional education and support from a health care professional. Specialty drugs offer treatment for serious, chronic, life-threatening diseases and are covered under medical benefits.

### DEFINITIONS

#### Applicable Drugs (Not All-Inclusive)

J-Code	Brand	Generic	Route of Administration	Notes
J0584	Crysvita	INJECTION BUROSUMAB-TWZA 1 MG	Subcutaneous	
J3241	Tepezza	teprotumumab-trbw	IV	
J7351	Durysta	bimatoprost implant	IV	
J1300	Soliris	Injection, eculizumab	IV	
J1823	Uplizna	INJECTION INEBILIZUMAB-CDON 1 MG	IV	
J1427	Viltepso	INJECTION VILTOLARSEN 10 MG	IV	
TBD	Nulibry	Injection, fosdenopterin	IV	

### POLICY

It is the policy of the Health Plan to maintain a prior authorization process that promotes appropriate utilization of specific drugs with potential for misuse or limited indications. This process involves a review using Food and Drug Administration (FDA) criteria to make a determination of Medical Necessity, and approval by the Medical Policy Committee.

The specialty drugs listed in this policy are subject to the prior authorization process.

### PROCEDURE

#### Initial Authorization Criteria:

*Must meet all of the criteria listed below:*

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- Must be prescribed for an FDA approved or compendia supported indication
- Must be used consistently with manufacturer’s prescribing information (e.g., contraindications, limitations, etc.)
- Must be prescribed at a dose within the manufacturer’s dosing guidelines (based on diagnosis, weight, etc.) listed in the FDA approved labeling
- Member must meet one of the following:
  - Be included within the patient population identified in the indication OR
  - Meet the eligibility criteria for the clinical studies

**Reauthorization Criteria:**

All prior authorization renewals are reviewed on an annual basis to determine the Medical Necessity for continuation of therapy. Authorization may be extended at 1-year intervals based upon chart documentation from the prescriber that the member’s condition has improved based upon the prescriber’s assessment while on therapy.

**Limitations:**

<b>Length of Authorization (if above criteria met)</b>	
Initial Authorization	Up to 1 year
Reauthorization	Same as initial

If the established criteria are not met, the request is referred to a Medical Director for review, if required for the plan and level of request.

**REFERENCES**

N/A

**REVIEW HISTORY**

<b>DESCRIPTION OF REVIEW / REVISION</b>	<b>DATE APPROVED</b>
Initial Review	3/22
Updated authorization durations to 1 year	2/23

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**Record Retention**

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