

Clinical Pharmacy Program Guidelines for Rayos

Program	Prior Authorization
Medication	Rayos (prednisone delayed-release tablet)
Markets in Scope	Arizona, California, Colorado, Hawaii, Maryland, Nevada, New Jersey, New York, New York EPP, Pennsylvania- CHIP, Rhode Island, South Carolina
Issue Date	9/2019
Pharmacy and Therapeutics Approval Date	9/2020
Effective Date	11/2020

1. Background:

Rayos (prednisone delayed-release tablet) is a corticosteroid indicated:

- as an anti-inflammatory or immunosuppressive agent for certain allergic, dermatologic, gastrointestinal, hematologic, ophthalmologic, nervous system, renal, respiratory, rheumatologic, specific infectious diseases or conditions and organ transplantation
- for the treatment of certain endocrine conditions
- for palliation of certain neoplastic conditions

2. Coverage Criteria:

<p>A. Rayos will be approved based on all of the following criteria:</p> <p style="margin-left: 40px;">1. <u>One</u> of the following:</p> <p style="margin-left: 80px;">(a) The requested drug must be used for an FDA-approved indication</p> <p style="text-align: center; margin-left: 40px;">-OR-</p> <p style="margin-left: 80px;">(b) The use of this drug is supported by information from the appropriate compendia of current literature.*</p> <p style="text-align: center; margin-left: 40px;">-AND-</p> <p style="margin-left: 40px;">2. The drug is being prescribed for a medically accepted indication that is recognized as a covered benefit by the applicable health plan’s program.</p> <p style="text-align: center; margin-left: 40px;">-AND-</p> <p style="margin-left: 40px;">3. Submission of medical records (e.g. chart notes, laboratory values) documenting an intolerance to generic prednisone tablets which is unable to</p>
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be resolved with attempts to minimize the adverse effects where appropriate.

-AND-

4. History of failure, contraindication, or intolerance to **two** of the following:

- Dexamethasone tablet/oral solution
- Hydrocortisone tablet
- Methylprednisolone tablet
- Prednisolone tablet/oral solution

Authorization will be issued for 12 months.

*Compendia of current literature: • American Hospital Formulary Service Drug Information • National Comprehensive Cancer Network Drugs and Biologics Compendium • Thomson Micromedex DrugDex • Clinical Pharmacology • United States Pharmacopoeia-National Formulary (USP-NF)

3. Additional Clinical Rules:

- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class.
- Supply limits may be in place.

4. References:

1. Rayos [package insert]. Lake Forest, IL: Horizon Pharma USA, Inc.; December 2019.

Program	Prior Authorization –Rayos
Change Control	
Date	Change
9/2019	New policy
9/2020	Annual review. Removed prednisolone oral syrup as a step therapy drug. Updated references.