

# Pharmacy clinical criteria

**Effective Feb. 15, 2026**, these are clinical criteria updates for UnitedHealthcare Community Plan of Texas CHIP, STAR, STAR Kids and STAR+PLUS plans.

Clinical criteria guidelines	Clinical criteria updates
<b>Allergen Extracts</b>	<ul style="list-style-type: none"> <li>• Odactra: added criteria</li> <li>• Odactra: updated minimum age to 5 years (from 12 years)</li> <li>• Palforzia: updated minimum age to 1 year (from 4 years)</li> </ul>
<b>Calcitonin Gene-Related Peptide Receptor (CGRP) Antagonists, Prophylaxis</b>	<ul style="list-style-type: none"> <li>• Ajovy: updated age check for patients 6 years and older with episodic migraine diagnosis</li> </ul>
<b>Cystic Fibrosis Agents</b>	<ul style="list-style-type: none"> <li>• Alyftrek: added criteria as approved by the Drug Utilization Review Board (DUR)</li> </ul>
<b>Cytokine and CAM Antagonists</b>	<ul style="list-style-type: none"> <li>• Legselvi: added criteria as approved by the DUR Board</li> <li>• Tremfya: added criteria to check for Crohn's disease</li> <li>• Rinvoq: added criteria to check for giant cell arteritis</li> </ul>
<b>Dipeptidyl Peptidase-4 (DPP-4) Inhibitors</b>	<ul style="list-style-type: none"> <li>• New prior authorization</li> </ul>
<b>Duchenne Muscular Dystrophy (DMD) Agents</b>	<ul style="list-style-type: none"> <li>• Emflaza: HHSC retired the prior authorization criteria and Emflaza Prior Authorization Request (HHS Form 1347). The Duchenne Muscular Dystrophy Agents criteria guide now includes Emflaza.</li> <li>• Duvyzat: added criteria as approved by the DUR Board</li> <li>• Changed the criteria guide's name from "Systemic Corticosteroids" to "Duchenne Muscular Dystrophy Agents"</li> </ul>
<b>Gaucher's Disease Agents</b>	<ul style="list-style-type: none"> <li>• Removed the negating diagnosis for pregnancy table (no longer a part of the criteria)</li> </ul>
<b>Glatiramer</b>	<ul style="list-style-type: none"> <li>• Copaxone: added to the Drugs Requiring PA section</li> </ul>
<b>Hereditary Angioedema Agents</b>	<ul style="list-style-type: none"> <li>• Kalbitor: removed from the Drugs Requiring PA section - Healthcare professional administered only</li> <li>• Dawnzera: added criteria as approved by the DUR Board</li> </ul>

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<b>Hyperlipidemia Agents</b>	<ul style="list-style-type: none"> <li>• Praluent: removed check for a PA form, updated question 2 to say "Deny" if the answer is "No", and removed the manual check for the diagnosis of HeFH</li> <li>• Repatha: removed check for a PA form</li> <li>• PCSK9 products: HHSC retired the prior authorization criteria and PCSK9 Inhibitors Prior Authorization Request (HHS Form 1355). The Hyperlipidemia Agents clinical criteria guide now includes PCSK9 products.</li> </ul>
<b>Imcivree</b>	<ul style="list-style-type: none"> <li>• Updated question 1 in the initial review criteria from 6 years to 2 years</li> <li>• Updated the approval duration to 365 days for initial approval for all indications</li> <li>• Added a check for end-stage renal disease (ESRD) to renewal criteria</li> </ul>
<b>Immunomodulators, Topical</b>	<ul style="list-style-type: none"> <li>• Anzupgo: added to the Opzelura Drugs Requiring PA section</li> <li>• Opzelura: updated age check to 2 years of age and older for atopic dermatitis indication</li> </ul>
<b>Monoclonal Antibody Agents</b>	<ul style="list-style-type: none"> <li>• Dupixent: added criteria for to check for bullous pemphigoid</li> <li>• Dupixent: added new indication of chronic spontaneous urticaria as approved by the DUR Board</li> <li>• Nucala: added new indication of chronic obstructive pulmonary disease and an eosinophilic phenotype as approved by the DUR Board</li> <li>• Airduo Digihaler: removed from the Supporting Tables section – products discontinued</li> </ul>
<b>Opioid Policy</b>	<ul style="list-style-type: none"> <li>• Added max days' supply check of 34 days for subsequent fills</li> </ul>
<b>Oxybate Products</b>	<ul style="list-style-type: none"> <li>• Lumryz: added to the Drugs Requiring PA section</li> </ul>
<b>Phosphodiesterase-5 (PDE-5) Inhibitors</b>	<ul style="list-style-type: none"> <li>• Ofsynvi: removed from the Drugs Requiring Prior Authorization section</li> <li>• Ofsynvi: refer to the Pulmonary Hypertension Agents criteria guide</li> </ul>
<b>Pulmonary Hypertension Agents</b>	<ul style="list-style-type: none"> <li>• Ofsynvi: added to Drugs Requiring Prior Authorization section</li> <li>• Winrevair and Yutrepia: added to the Drugs Requiring PA section</li> <li>• Ventavis: removed from the Drugs Requiring PA section and supporting tables – discontinued</li> <li>• Remodulin: added to the Injectable PH Drugs Requiring PA section</li> </ul>
<b>Topical Antifungals for Onychomycosis</b>	<ul style="list-style-type: none"> <li>• New prior authorization</li> </ul>
<b>Transthyretin Agents</b>	<ul style="list-style-type: none"> <li>• Tegsedi: removed criteria – product discontinued</li> <li>• Wainua: added criteria as approved by the DUR Board</li> </ul>
<b>Xifaxan</b>	<ul style="list-style-type: none"> <li>• Added levofloxacin to question 3 and Table 3 for Xifaxan 200 mg</li> </ul>