

The following changes will be effective on **February 1, 2025**, unless otherwise specified and apply to the following plan:

### Yamhill Community Care (Medicaid)

#### Formulary Changes

Drug Name	Formulary Status	Policy Name
<b>Febuxostat (Uloric) Tablet</b>	<ul style="list-style-type: none"> <li>Medicaid: Add generic to formulary</li> </ul>	N/A
<b>Chenodiol (Chenodal) Tablet</b>	Remove from Medicaid formulary <b>Effective: 03/01/2025</b>	Chenodal
<b>Cholic acid (Cholbam) Capsule</b>	Remove from Medicaid formulary <b>Effective: 03/01/2025</b>	Medications For Rare Indications
<b>Teduglutide (Gattex) Kit</b>	Remove from Medicaid formularies	Gattex
<b>Daprodustat (Jesduvroq) Tablet</b>	Remove from Medicaid formularies	Jesduvroq, Vafseo
<b>Obeticholic acid (Ocaliva) Tablet</b>	Remove from Medicaid formulary	Primary Biliary Cholangitis Agents
<b>Granisetron (Sancuso) Patch TDWK</b>	Remove from Medicaid formulary	N/A

<b>Budesonide (Uceris) 9 mg Tab DR/ER</b>	<ul style="list-style-type: none"> <li>Medicaid: Remove from formulary and add Quantity Limit (one tablet per day) <b>Effective: 03/01/2025</b></li> </ul>	Uceris
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## Medical Policy Changes

### Coverage Criteria Changes

Drug/Policy Name(s)	Plans Affected	Summary of Change
<b>Anti-Cancer Medications – Self-Administered</b>	<input checked="" type="checkbox"/> Medicaid	Will require trial of imatinib before coverage of nilotinib (Tasigna®) and dasatinib (Sprycel®) will be authorized. This will apply to new starts only.
<b>Acute Hereditary Angioedema Therapy</b>	<input checked="" type="checkbox"/> Medicaid	Updated age restrictions language to require age be appropriate based on FDA approved indication. Updated quantity limit for icatibant to allow treatment for two exacerbations per month. Per package insert, may administer up to three doses per 24 hours.
<b>Adakveo</b>	<input checked="" type="checkbox"/> Medicaid	Removed Oxbryta® under exclusion since the drug has been withdrawn from the market. Added Endari® under exclusion criteria to prevent combination use.
<b>Alinia</b>	<input checked="" type="checkbox"/> Medicaid	Updated oral suspension quantity limit to allow for three days of treatment per package insert.
<b>Chenodal</b>	<input checked="" type="checkbox"/> Medicaid	Added requirement for radiolucent stones in well-opacifying gallbladders, and clarifide when patients are considered not a candidate for surgery.

<b>Drug/Policy Name(s)</b>	<b>Plans Affected</b>	<b>Summary of Change</b>
<b>Constipation Agents – Medicaid</b>	<input checked="" type="checkbox"/> Medicaid	Added criteria for reauthorization criteria, updated ICD-10 code list of non-covered diagnosis codes to include functional constipation as they are also considered unfunded diagnoses.
<b>Gene Therapies for Hemoglobin Disorders</b>	<input checked="" type="checkbox"/> Medicaid	Added note that additional genotypes will be considered on a case-by-case basis based on disease severity for sickle cell disease to meet compliance for value based agreement.
<b>Givlaari</b>	<input checked="" type="checkbox"/> Medicaid	Updated active disease definition to include four or more porphyria attacks within a year (in addition to two or more within the past six months). This aligns with expert opinion statement from American Gastroenterological Association. Added for reauthorization that dosing must align with FDA-labeling.
<b>Hemlibra</b>	<input checked="" type="checkbox"/> Medicaid	Added criteria requiring the dose and frequency align with FDA labeling.
<b>Hepatitis C - Direct Acting Antivirals</b>	<input checked="" type="checkbox"/> Medicaid	Allow coverage of generic Epclusa in solid organ transplant setting per AASLD guideline.
<b>Jesduvroq, Vafseo</b>	<input checked="" type="checkbox"/> Medicaid	Updated prescriber restrictions to allow hematologist. Updated duration of approval to align with Erythropoietin Stimulating Agents clinical policy.
<b>Livtency</b>	<input checked="" type="checkbox"/> Medicaid	Updated criteria to require failure of one antiviral or intolerance/contraindication to all other listed antivirals.
<b>Lotronex</b>	<input checked="" type="checkbox"/> Medicaid	Removed information on REMS program in prescriber restrictions and position statement as this is no longer required, increased initial authorization duration to 12 months, removed requirement that patient is female due to low risk of inappropriate utilization and low likelihood of males continuing on therapy if they are approved due to decreased efficacy in this population.

Drug/Policy Name(s)	Plans Affected	Summary of Change
<b>Medications For Rare Indications</b>	<input checked="" type="checkbox"/> Medicaid	For Cerdelga, add requirement for metabolic status of poor, intermediate, or extensive 2D6 metabolizer. For Galafold, updated diagnosis requirement to an amenable galactosidase alpha (GLA) gene variant. For Sohonos, required initial clinical scores. For Xolremdi, required initial labs.
<b>Prevyomis</b>	<input checked="" type="checkbox"/> Medicaid	Updated age restriction as medication is now approved down to those 6 months of age for hematopoietic stem cell transplantation (HSCT) and 12 years of age for kidney transplant recipients. Clarified that coverage requests for HSCT greater than 100 days post transplantation requires documentation the member is at high risk for late cytomegalovirus infection.
<b>Primary Biliary Cholangitis Agents</b>	<input checked="" type="checkbox"/> Medicaid	Updated initial auth from four to six months to allow more time to assess response.
<b>Reblozyl, Rytelo</b>	<input checked="" type="checkbox"/> Medicaid	Clarified definition of transfusion-dependent anemia for beta-thalassemia.
<b>Tavneos</b>	<input checked="" type="checkbox"/> Medicaid	Coverage duration clarified.
<b>Thrombocytopenia Medications</b>	<input checked="" type="checkbox"/> Medicaid	For Immune Thrombocytopenia (ITP), removed rituximab as trial/failure option; For Severe aplastic anemia (AA), added requirement for combination or previous use of standard immunosuppressive therapy; For Chronic Liver Disease, removed requirement for when to start therapy; For continuation of ITP and AA, remove requirement for attestation of medical necessity; Added quantity limits to Doptelet and Promacta. <b>Effective 03/01/2025</b>
<b>Uceris</b>	<input checked="" type="checkbox"/> Medicaid	Removed tablet from Medicaid formulary and added quantity limit for tablet. <b>Effective 03/01/2025</b>

### Retired Medical Policies

- **Altuviiiio**
- **Oxbryta – medication is no longer available**
- **Serotonin Antagonists Step Therapy Policy**

**New Drugs:**

<b>Drug Name</b>	<b>Recommendations</b>	<b>Policy Name</b>
<b>Donanemab-azbt (Kisunla) Vial</b>	<ul style="list-style-type: none"> <li>• Medicaid: Medical Benefit, Prior Authorization</li> </ul>	Anti-Amyloid Monoclonal Antibodies
<b>Xanomeline tart-trospium chlor (Cobenfy) Capsule</b>	<ul style="list-style-type: none"> <li>• Medicaid: Non-Formulary</li> </ul>	<ul style="list-style-type: none"> <li>• Commercial: Antipsychotics</li> <li>• Medicaid: N/A</li> </ul>
<b>Afamitresgene autoleucel (Tecelra) Plast. Bag</b>	<ul style="list-style-type: none"> <li>• Medicaid: Medical Benefit, Prior Authorization, Quantity Limit (1 dose per lifetime)</li> </ul>	T-Cell Therapy
<b>Arimoclomol citrate (Miplyffa) Capsule</b>	<ul style="list-style-type: none"> <li>• Medicaid: Non- Formulary, Prior Authorization, Quantity Limit (3 capsules per day)</li> </ul>	Medications for Rare Indications
<b>Bexagliflozin (Brenzavvy) Tablet</b>	<ul style="list-style-type: none"> <li>• Commercial/Medicaid: Non-Formulary</li> </ul>	N/A
<b>Lazertinib mesylate (Lazcluze) Tablet</b>	<ul style="list-style-type: none"> <li>• Medicaid: Non- Formulary, Prior Authorization, Quantity Limit (2 tablets per day for 80 mg; 1 tablet per day for 240 mg)</li> </ul>	Anti-Cancer Medications – Self-administered
<b>Lebrikizumab-lbkz (Ebglyss Pen) Pen Injctr</b>	<ul style="list-style-type: none"> <li>• Medicaid: Non-Formulary, Prior Authorization, Quantity Limit (2 mL per 28 days)</li> </ul>	Interleukin-13 Inhibitors
<b>Seladelpar lysine (Livdelzi) Capsule</b>	<ul style="list-style-type: none"> <li>• Medicaid: Non-Formulary, Prior Authorization, Quantity Limit (1 capsule per day)</li> </ul>	Primary Biliary Cholangitis Agents
<b>Tislelizumab-jsgr (Tevimbra) Vial</b>	<ul style="list-style-type: none"> <li>• Medicaid: Medical Benefit, Prior Authorization</li> </ul>	Anti-Cancer Medications Policy – Medical Benefit

<p><b>Vorasidenib citrate (Vorinigo) Tablet</b></p>	<ul style="list-style-type: none"> <li>Medicaid: Formulary, Prior Authorization, Quantity Limit (10 mg tablets: 60 tablets per 30 days; 40 mg tablets: 30 tablets per 30 days)</li> </ul>	<p>Anti-Cancer Medications – Self-Administered</p>
<p><b>Palopegteriparatide (Yorvipath) Pen Injector</b></p>	<ul style="list-style-type: none"> <li>Medicaid: Formulary, Prior Authorization, Quantity Limit (2 pens per 28 days)</li> </ul>	<p>Yorvipath</p>