

**PROPOSED CRITERIA:**

**Type:**  New Criteria       Revised Existing Criteria       PA Group drug list change, criteria remains the same

**Applies to:**       New Starts Only       New and existing users -- member notification is required when a drug is removed from formulary, but not when criteria changes for NF drugs; consult PHC Policy MCRP4064, Continuation of Prescription Drugs.

**PA Group Name:** *Antineoplastic Drugs*

**PA Group Includes:** *Antineoplastic Agents not otherwise having drug-specific PA Criteria*

**Recommended Criteria:**

<b>Covered Uses</b>	<ul style="list-style-type: none"> <li>• Case-Specific.</li> <li>• FDA approved indications.</li> <li>• Off-Label indications: medically accepted indications are defined using the following standard reference compendia, under the Centers for Medicare and Medicaid Services guidance: American Hospital Formulary Service-Drug Information (AHFS-DI), Truven Health Analytics Micromedex DrugDeX (DrugDex), National Comprehensive Cancer Network (NCCN) Drugs and Biologics Compendium (as indicated by a category 1, 2A, or 2B), Wolters Kluwer Lexi-Drugs, Elsevier/Gold Standard Clinical Pharmacology and/or positive results from two peer-reviewed published medical studies.</li> </ul>
<b>Reasons for Exclusion</b>	Uses without supporting evidence for the stated indication (experimental).
<b>Required Medical Documentation</b>	TAR must include accurate diagnosis as provided by PRESCRIBER and include all necessary/relevant clinical documentation to support medical justification (e.g. clinic notes, treatment history including prior regimen(s), lab reports, specialist consults, imaging reports, etc).
<b>Age Restrictions</b>	
<b>Prescriber Restriction</b>	Prescribed by oncologist or hematologist, or with specialist consult/recommendation.
<b>Coverage Duration</b>	Initial: When applicable, 14 day supply per fill, during the first two months of therapy Renewal: 6 month intervals
<b>Other Requirements</b>	<p>Renewal: requires clinical documentation demonstrating that the patient is demonstrating a positive response to the requested therapy (as evidenced by an improvement in the condition being treated without adverse effects causing treatment interruption), and any current, most updated assessment/treatment plan for this patient.</p> <p><input checked="" type="checkbox"/> LD/SP: Limited to dispensing by AllianceRx/Walgreens <i>when applicable</i></p> <p><input checked="" type="checkbox"/> May be limited to a 14-15 day supply for the first 2 months of treatment, until dose-stable, when product packaging allows partial-package dispensing.</p>
<input type="checkbox"/> Approved <input type="checkbox"/> Approved as Modified:	

**NCCN Categories of Evidence and Consensus**

**Category 1:** Based upon high-level evidence, there is uniform NCCN consensus that the intervention is appropriate.

**Category 2A:** Based upon lower-level evidence, there is uniform NCCN consensus that the intervention is appropriate.

**Category 2B:** Based upon lower-level evidence, there is NCCN consensus that the intervention is appropriate.

**Category 3:** Based upon any level of evidence, there is major NCCN disagreement that the intervention is appropriate.