

Clinical Pharmacy Program Guidelines for Qinlock

Program	Prior Authorization
Medication	Qinlock™ (ripretinib)
Markets in Scope	Arizona, California, Hawaii, Nevada, Maryland, New Jersey, New York, New York EPP, Pennsylvania- CHIP, Rhode Island, South Carolina
Issue Date	7/2020
Pharmacy and Therapeutics Approval Date	7/2020
Effective Date	9/2020

1. Background:

Qinlock (ripretinib) is a kinase inhibitor indicated for the treatment of adult patients with advanced gastrointestinal stromal tumor (GIST) who have received prior treatment with 3 or more kinase inhibitors, including imatinib.

The National Cancer Comprehensive Network (NCCN) recommends Qinlock as the preferred fourth-line therapy for unresectable or metastatic disease progressive after single agent therapy with imatinib, sunitinib, and regorafenib.

2. Coverage Criteria:

<p><u>A. Gastrointestinal Stromal Tumor (GIST)</u></p> <p>1. <u>Initial Authorization</u></p> <p style="padding-left: 40px;">a. Qinlock will be approved based on <u>all</u> of the following criteria:</p> <p style="padding-left: 80px;">(1) Diagnosis of gastrointestinal stromal tumor (GIST)</p> <p style="text-align: center;">-AND-</p> <p style="padding-left: 80px;">(2) Disease is <u>one</u> of the following:</p> <p style="padding-left: 120px;">(a) Advanced (b) Metastatic (c) Unresectable</p> <p style="text-align: center;">-AND-</p>
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(3) History of failure to **all** of the following:

- (a) imatinib (Gleevec)
- (b) sunitinib (Sutent)
- (c) regorafenib (Stivarga)

Authorization will be issued for 12 months.

2. Reauthorization

a. **Qinlock** will be approved based on the following criterion:

- (1) Patient does not show evidence of progressive disease while on Qinlock therapy

Authorization will be issued for 12 months.

B. NCCN Recommended Regimens

1. Initial Authorization

a. **Qinlock** will be approved for uses not outlined above if supported by The National Comprehensive Cancer Network (NCCN) Drugs and Biologics Compendium.

Authorization will be issued for 12 months.

2. Reauthorization

a. **Qinlock** will be approved based on the following criterion:

- (1) Documentation of positive clinical response to Qinlock therapy

Authorization will be issued for 12 months.

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. Qinlock [package insert]. Waltham, MA: Deciphera Pharmaceuticals, LLC; May 2020.
2. The NCCN Drugs and Biologics Compendium (NCCN Compendium™). Available at www.nccn.org. Accessed May 22, 2020

Program	Program type – Prior Authorization
Change Control	
7/2020	New program.