

### UnitedHealthcare Pharmacy Clinical Pharmacy Programs

Program Number	2024 P 1092-12
Program	Prior Authorization/Notification
Medication	Selzentry <sup>®</sup> (maraviroc)
P&T Approval Date	1/2012, 2/2013, 11/2013, 2/2015, 2/2016, 2/2017, 2/2018, 2/2019,
	2/2020, 2/2021, 1/2022, 1/2023, 1/2024
Effective Date	4/1/2024

## 1. Background:

Selzentry<sup>®</sup> (maraviroc) is a CCR5 co-receptor antagonist indicated in combination with other antiretroviral agents for the treatment of *only* CCR5-tropic human immunodeficiency virus type 1 (HIV-1) infection in adults and pediatric patients weighing at least 2 kg. Selzentry is not recommended in patients with dual/mixed- or CXCR4-tropic HIV-1. Tropism testing with a highly sensitive tropism assay is required for the appropriate use of Selzentry.<sup>1</sup>

Members will be required to meet the coverage criteria below.

## 2. Coverage Criteria<sup>a</sup>:

# A. <u>Selzentry</u>

- 1. Selzentry will be approved based on <u>both</u> of the following criteria:
  - a. Patient has CCR5-tropic HIV-1 infection as confirmed by a highly sensitive tropism assay

#### -AND-

b. Patient is currently taking or will be prescribed an optimized background antiretroviral therapy regimen

#### Authorization will be issued for 12 months.

<sup>a</sup> State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

#### 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. Selzentry [Package Insert]. Research Triangle Park, NC: ViiV Healthcare; September 2022.

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Change Control	
11/2013	Annual Review - Removed reauthorization criteria. Clinical intent is to validate tropism assay testing at time of therapy initiation. Updated additional clinical rules. Updated references. Added change control table.
2/2015	Removed safety information in background section and added updated label language. Updated references.
2/2016	Annual review. Updated background section to reflect most current label. Removed reference to tropism testing from the DHHS treatment guidelines. Revised duration of authorization.
2/2017	Annual Review. Updated background information to reflect most current label. Updated reference.
2/2018	Annual review. No changes to coverage criteria.
12/2018	Administrative change to add statement regarding use of automated processes.
2/2019	Annual review. No changes to coverage criteria.
2/2020	Annual review. No changes to coverage criteria.
2/2021	Annual review. No changes to coverage criteria.
1/2022	Annual review with no changes to coverage criteria. Updated background and reference.
1/2023	Annual review with no changes to coverage criteria. Added state mandate and updated references.
1/2024	Annual review. Revised duration of authorization.