



**Scemblix® (asciminib)**  
Effective 09/01/2022

<b>Plan</b>	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	<b>Program Type</b>	<input checked="" type="checkbox"/> Prior Authorization <input checked="" type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
<b>Benefit</b>	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit (NLX)		
<b>Specialty Limitations</b>	This medication has been designated specialty and must be filled at a contracted specialty pharmacy.		
<b>Contact Information</b>	<b>Specialty Medications</b>		
	All Plans	Phone: 866-814-5506	Fax: 866-249-6155
	<b>Non-Specialty Medications</b>		
	MassHealth	Phone: 877-433-7643	Fax: 866-255-7569
	Commercial	Phone: 800-294-5979	Fax: 888-836-0730
	Exchange	Phone: 855-582-2022	Fax: 855-245-2134
	<b>Medical Specialty Medications (NLX)</b>		
	All Plans	Phone: 844-345-2803	Fax: 844-851-0882
<b>Exceptions</b>	N/A		

**Overview**

Scemblix is FDA approved for:

1. Adult patients with Philadelphia chromosome-positive chronic myeloid leukemia in chronic phase (Ph+ CML-CP), previously treated with two or more tyrosine kinase inhibitors (TKIs)
2. Adult patients with Philadelphia chromosome-positive chronic myeloid leukemia in chronic phase (Ph+ CML-CP) with the T315I mutation

**Coverage Guidelines**

Authorization may be reviewed for members new to AllWays Health Partners who are currently receiving treatment with Scemblix, excluding when the product is obtained as samples or via manufacturer’s patient assistance programs

**OR**

Authorization may be granted for members when all the following criteria are met, and documentation is provided:

1. Member has a diagnosis of Philadelphia chromosome (Ph+) chronic myeloid leukemia (CML) in chronic phase
2. Member has ONE of the following:
  - a. Member has T315I mutation positive CML
  - b. Provider attestation that member has been previously treated with at least two kinase inhibitors (e.g., bosutinib, dasatinib, imatinib, nilotinib)

**Continuation of Therapy**

Reauthorizations requires physician attestation of continuation of therapy and no evidence of unacceptable toxicity or disease progression while on the current regimen.



**Limitations**

1. Initial approvals and reauthorizations will be granted for 12 months
2. The following quantity limits apply:

Scemblix 40mg	300 tablets per 30 days
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**References**

1. Scemblix [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; October 2021.

**Review History**

06/22/2022 – Created and reviewed for June P&T; Effective 09/01/2022

**Disclaimer**

AllWays Health Partners complies with applicable federal civil rights laws and does not discriminate or exclude people on the basis of race, color, national origin, age, disability, or sex.