

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use SCSEMBLIX® safely and effectively. See full prescribing information for SCSEMBLIX.

SCSEMBLIX (asciminib) tablets, for oral use
Initial U.S. Approval: 2021

INDICATIONS AND USAGE

SCSEMBLIX is a kinase inhibitor indicated for the treatment of adult patients with:

- Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic phase (CP), previously treated with two or more tyrosine kinase inhibitors (TKIs). (1)
This indication is approved under accelerated approval based on major molecular response (MMR). Continued approval for this indication may be contingent upon verification and description of clinical benefit in a confirmatory trial(s).
- Ph+ CML in CP with the T315I mutation. (1)

DOSAGE AND ADMINISTRATION

- Recommended Dosage in Ph+ CML in CP:** 80 mg orally once daily or 40 mg twice daily. (2.1)
- Recommended Dosage in Ph+ CML in CP with the T315I Mutation:** 200 mg orally twice daily. (2.2)
- Avoid food for at least 2 hours before and 1 hour after taking SCSEMBLIX. (2.5)
- Swallow tablets whole. Do not break, crush, or chew the tablets. (2.5)

DOSAGE FORMS AND STRENGTHS

- Film-coated tablets: 20 mg and 40 mg (3)

CONTRAINDICATIONS

None. (4)

WARNINGS AND PRECAUTIONS

- Myelosuppression:** Severe thrombocytopenia and neutropenia events may occur. Monitor complete blood counts regularly during therapy and manage by treatment interruption or dose reduction. (2.4, 5.1)
- Pancreatic Toxicity:** Monitor serum lipase and amylase. Interrupt, then resume at reduced dose or discontinue SCSEMBLIX based on severity. Evaluate for pancreatitis when lipase elevation is accompanied by abdominal symptoms. (2.4, 5.2)
- Hypertension:** Monitor blood pressure and manage hypertension as clinically indicated. Interrupt, dose reduce, or stop SCSEMBLIX if hypertension is not medically controlled. (2.4, 5.3)

- Hypersensitivity:** May cause hypersensitivity reactions. Monitor patients for signs and symptoms and initiate appropriate treatment as clinically indicated. (5.4)
- Cardiovascular Toxicity:** Cardiovascular toxicity may occur. Monitor patients with history of cardiovascular risk factors for cardiovascular signs and symptoms. Initiate appropriate treatment as clinically indicated. (5.5)
- Embryo-Fetal Toxicity:** Can cause fetal harm. Advise females of reproductive potential of the potential risk to a fetus and to use effective contraception. (5.6, 8.1, 8.3)

ADVERSE REACTIONS

Most common adverse reactions ($\geq 20\%$) are upper respiratory tract infections, musculoskeletal pain, fatigue, nausea, rash, and diarrhea. (6.1)
Most common laboratory abnormalities ($\geq 20\%$) are platelet count decreased, triglycerides increased, neutrophil count decreased, hemoglobin decreased, creatine kinase increased, alanine aminotransferase increased, lipase increased, and amylase increased. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Novartis Pharmaceuticals Corporation at 1-888-669-6682 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Strong CYP3A4 Inhibitors:** Closely monitor for adverse reactions during concomitant use of SCSEMBLIX at 200 mg twice daily. (7.1)
- Itraconazole Oral Solution Containing Hydroxypropyl- β -cyclodextrin:** Avoid concomitant use of SCSEMBLIX at all recommended doses. (7.1)
- Certain Substrates of CYP3A4:** Closely monitor for adverse reactions during concomitant use of SCSEMBLIX at 80 mg total daily dose. Avoid use of SCSEMBLIX at 200 mg twice daily. (7.2)
- Substrates of CYP2C9:** Avoid concomitant use of SCSEMBLIX at all recommended doses.
 - 80 mg total daily dose:** If unavoidable, reduce the CYP2C9 substrate dosage as necessary. (7.2)
 - 200 mg twice daily:** If unavoidable, consider alternative therapy with non-CYP2C9 substrate. (7.2)
- Certain P-gp Substrates:** Closely monitor for adverse reactions during concomitant use of SCSEMBLIX at all recommended doses. (7.2)

USE IN SPECIFIC POPULATIONS

Lactation: Advise not to breastfeed. (8.2)

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

SCEMBLIX is indicated for the treatment of adult patients with:

- Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic phase (CP), previously treated with two or more tyrosine kinase inhibitors (TKIs).

This indication is approved under accelerated approval based on major molecular response (MMR) [see *Clinical Studies (14.1)*]. Continued approval for this indication may be contingent upon verification and description of clinical benefit in a confirmatory trial(s).

- Ph+ CML in CP with the T315I mutation.

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage in Patients with Ph+ CML-CP, Previously Treated with Two or More TKIs

The recommended dose of SCEMBLIX is 80 mg taken orally once daily at approximately the same time each day or 40 mg twice daily at approximately 12-hour intervals. The recommended dose of SCEMBLIX is taken orally without food. Avoid food consumption for at least 2 hours before and 1 hour after taking SCEMBLIX [see *Clinical Pharmacology (12.2)*].

Continue treatment with SCEMBLIX as long as clinical benefit is observed or until unacceptable toxicity occurs.

2.2 Recommended Dosage in Patients with Ph+ CML-CP with the T315I Mutation

The recommended dose of SCEMBLIX is 200 mg taken orally twice daily at approximately 12-hour intervals. The recommended dose of SCEMBLIX is taken orally without food. Avoid food consumption for at least 2 hours before and 1 hour after taking SCEMBLIX [see *Clinical Pharmacology (12.2)*].

2.3 Missed Dose

Once Daily Dosage Regimen: If a SCEMBLIX dose is missed by more than approximately 12 hours, skip the dose and take the next dose as scheduled.

Twice Daily Dosage Regimens: If a SCEMBLIX dose is missed by more than approximately 6 hours, skip the dose and take the next dose as scheduled.

2.4 Dosage Modifications

Dosage Modifications for Patients with Ph+ CML-CP, Previously Treated with Two or More TKIs

For the management of adverse reactions, reduce the SCEMBLIX dose as described in Table 1.

Dosage Modifications for Patients with Ph+ CML-CP with the T315I Mutation

For the management of adverse reactions, reduce the SCEMBLIX dose as described in Table 1.

Table 1: Recommended Dosage Reductions for SCEMBLIX for Adverse Reactions		
Dosage Reduction	Dosage for Patients with CP-CML, Previously Treated with Two or More TKIs	Dosage for Patients with Ph+ CML-CP with the T315I Mutation
First	<ul style="list-style-type: none">40 mg once daily OR <ul style="list-style-type: none">20 mg twice daily	160 mg twice daily
Subsequent Reduction	Permanently discontinue SCEMBLIX in patients unable to tolerate 40 mg once daily OR 20 mg twice daily.	Permanently discontinue SCEMBLIX in patients unable to tolerate 160 mg twice daily.

The recommended dosage modifications for the management of selected adverse reactions is shown in Table 2.

Table 2: SCEMBLIX Dosage Modification for the Management of Adverse Reactions

Adverse Reaction	Dosage Modification
Thrombocytopenia and/or neutropenia [see <i>Warnings and Precautions (5.1)</i>]	
ANC less than $1.0 \times 10^9/L$ and/or PLT less than $50 \times 10^9/L$	<p>Withhold SCEMBLIX until resolved to ANC greater than or equal to $1 \times 10^9/L$ and/or PLT greater than or equal to $50 \times 10^9/L$.</p> <p>If resolved:</p> <ul style="list-style-type: none"> • Within 2 weeks: resume SCEMBLIX at starting dose. • After more than 2 weeks: resume SCEMBLIX at reduced dose. <p>For recurrent severe thrombocytopenia and/or neutropenia, withhold SCEMBLIX until resolved to ANC greater than or equal to $1 \times 10^9/L$ and PLT greater than or equal to $50 \times 10^9/L$, then resume at reduced dose.</p>
Asymptomatic amylase and/or lipase elevation [see <i>Warnings and Precautions (5.2)</i>]	
Elevation greater than $2.0 \times ULN$	<p>Withhold SCEMBLIX until resolved to less than $1.5 \times ULN$.</p> <p>If resolved:</p> <ul style="list-style-type: none"> • Resume SCEMBLIX at reduced dose. If events reoccur at reduced dose, permanently discontinue SCEMBLIX. <p>If not resolved:</p> <ul style="list-style-type: none"> • Permanently discontinue SCEMBLIX. Perform diagnostic tests to exclude pancreatitis.
Non-hematologic adverse reactions [see <i>Warnings and Precautions (5.3, 5.4, 5.5)</i>]	
Grade 3 ¹ or higher	<p>Withhold SCEMBLIX until recovery to Grade 1 or less.</p> <p>If resolved:</p> <ul style="list-style-type: none"> • Resume SCEMBLIX at reduced dose. <p>If not resolved:</p> <ul style="list-style-type: none"> • Permanently discontinue SCEMBLIX.
<p>Abbreviations: ANC, absolute neutrophil count; PLT, platelets; ULN, upper limit of normal. ¹Based on Common Terminology Criteria for Adverse Events (CTCAE) v 4.03.</p>	

2.5 Administration

Advise patients to swallow SCEMBLIX tablets whole. Do not break, crush, or chew the tablets.

3 DOSAGE FORMS AND STRENGTHS

- 20 mg film-coated tablets: pale yellow, unscored, round, biconvex, with beveled edges, film-coated tablet debossed with “20” on one side and “Novartis logo” on the other side.
- 40 mg film-coated tablets: violet white, unscored, round, biconvex, with beveled edges, film-coated tablet debossed with “40” on one side and “Novartis logo” on the other side.

4 CONTRAINDICATIONS

None.

5 WARNINGS AND PRECAUTIONS

5.1 Myelosuppression

Thrombocytopenia, neutropenia, and anemia have occurred in patients receiving SCEMBLIX. Thrombocytopenia occurred in 98 of 356 (28%) patients receiving SCEMBLIX, with Grade 3 or 4 thrombocytopenia reported in 24 (7%) and 42 (12%) of patients, respectively. Among the patients with Grade 3 or 4 thrombocytopenia, median time to first occurrence of events was 6 weeks (range, 0.1 to 64 weeks). Of the 98 patients with thrombocytopenia, 7 (2%) patients permanently discontinued SCEMBLIX, while SCEMBLIX was temporarily withheld in 45 (13%) patients due to the adverse reaction.

Neutropenia occurred in 69 (19%) patients receiving SCEMBLIX, with Grade 3 and 4 neutropenia reported in 27 (8%) and 29 (8%) patients, respectively. Among the patients with Grade 3 or 4 neutropenia, median time to first occurrence of events was 6 weeks (range, 0.1 to 180 weeks) Of the 69 patients with neutropenia, 4 (1.1%) patients permanently discontinued SCEMBLIX, while SCEMBLIX was temporarily withheld in 34 (10%) patients due to the adverse reaction.

Anemia occurred in 45 (13%) patients receiving SCEMBLIX, with Grade 3 anemia occurring in 19 (5%) patients. Among the patients with Grade 3 or 4 anemia, median time to first occurrence of events was 30 weeks (range, 0.4 to 207 weeks). Of the 45 patients with anemia, SCEMBLIX was temporarily withheld in 2 (0.6%) patients due to the adverse reaction [see Adverse Reactions (6.1)].

Perform complete blood counts every two weeks for the first 3 months of treatment and monthly thereafter or as clinically indicated. Monitor patients for signs and symptoms of myelosuppression.

Based on the severity of thrombocytopenia and/or neutropenia, reduce dose, temporarily withhold, or permanently discontinue SCEMBLIX [see Dosage and Administration (2.4)].

5.2 Pancreatic Toxicity

Pancreatitis occurred in 9 of 356 (2.5%) patients receiving SCEMBLIX, with Grade 3 pancreatitis occurring in 4 (1.1%) patients. All cases of pancreatitis occurred in the Phase I study (X2101). Of the 9 patients with pancreatitis, two (0.6%) patients permanently discontinued SCEMBLIX, while SCEMBLIX was temporarily withheld in 4 (1.1%) patients due to the adverse reaction. Asymptomatic elevation of serum lipase and amylase occurred in 76 of 356 (21%) patients receiving SCEMBLIX, with Grade 3 and Grade 4 pancreatic enzyme elevations occurring in 36 (10%) and 8 (2.2%) patients, respectively. Of the 76 patients with pancreatic enzymes elevated, SCEMBLIX was permanently discontinued in 7 (2%) patients due to the adverse reaction [see Adverse Reactions (6.1)].

Assess serum lipase and amylase levels monthly during treatment with SCEMBLIX, or as clinically indicated. Monitor patients for signs and symptoms of pancreatic toxicity. Perform more frequent monitoring in patients with a history of pancreatitis. If lipase and amylase elevation are accompanied by abdominal symptoms, temporarily withhold SCEMBLIX and consider appropriate diagnostic tests to exclude pancreatitis [see Dosage and Administration (2.4)].

Based on the severity of lipase and amylase elevation, reduce dose, temporarily withhold, or permanently discontinue SCEMBLIX [see Dosage and Administration (2.4)].

5.3 Hypertension

Hypertension occurred in 66 of 356 (19%) patients receiving SCEMBLIX, with Grade 3 or 4 hypertension reported in 31 (9%) and 1 (0.3%) patients, respectively. Among the patients with Grade 3 or 4 hypertension, median time to first occurrence was 14 weeks (range, 0.1 to 156 weeks). Of the 66 patients with hypertension, SCEMBLIX was temporarily withheld in 3 (0.8%) patients due to the adverse reaction [see Adverse Reactions (6.1)].

Monitor and manage hypertension using standard antihypertensive therapy during treatment with SCEMBLIX as clinically indicated; for Grade 3 or higher hypertension, temporarily withhold, reduce dose, or permanently discontinue SCEMBLIX depending on persistence of hypertension [see Dosage and Administration (2.4)].

5.4 Hypersensitivity

Hypersensitivity occurred in 113 of 356 (32%) patients receiving SCEMBLIX, with Grade 3 or 4 hypersensitivity reported in 6 (1.7%) patients [see Adverse Reactions (6.1)]. Reactions included rash, edema, and bronchospasm. Monitor

patients for signs and symptoms of hypersensitivity and initiate appropriate treatment as clinically indicated; for Grade 3 or higher hypersensitivity, temporarily withhold, reduce dose, or permanently discontinue depending on SCSEMBLIX depending on persistence of hypersensitivity [see *Dosage and Administration* (2.4)].

5.5 Cardiovascular Toxicity

Cardiovascular toxicity (including ischemic cardiac and CNS conditions, arterial thrombotic and embolic conditions) and cardiac failure occurred in 46 (13%) and in 8 (2.2%) of 356 patients receiving SCSEMBLIX, respectively [see *Adverse Reactions* (6.1)]. Grade 3 cardiovascular toxicity was reported in 12 (3.4%) patients, while grade 3 cardiac failure was observed in 4 (1.1%) patients. Grade 4 cardiovascular toxicity occurred in 2 (0.6%) patients, with fatalities occurring in 3 (0.8%) patients. Permanent discontinuation of SCSEMBLIX occurred in 3 (0.8%) patients due to cardiovascular toxicity and in 1 (0.3%) patient due to cardiac failure, respectively. Cardiovascular toxicity occurred in patients with pre-existing cardiovascular conditions or risk factors, and/or prior exposure to multiple TKIs.

Arrhythmia, including QTc prolongation, occurred in 23 of 356 (7%) patients receiving SCSEMBLIX, with Grade 3 arrhythmia reported in 7 (2%) patients. QTc prolongation occurred in 3 of 356 (0.8%) patients receiving SCSEMBLIX, with Grade 3 QTc prolongation reported in 1 (0.3%) patient [see *Adverse Reactions* (6.1)].

Monitor patients with history of cardiovascular risk factors for cardiovascular signs and symptoms. Initiate appropriate treatment as clinically indicated; for Grade 3 or higher cardiovascular toxicity, temporarily withhold, reduce dose, or permanently discontinue SCSEMBLIX depending on persistence of cardiovascular toxicity [see *Dosage and Administration* (2.4)].

5.6 Embryo-Fetal Toxicity

Based on findings from animal studies and its mechanism of action, SCSEMBLIX can cause fetal harm when administered to a pregnant woman. In animal reproduction studies, administration of asciminib to pregnant rats and rabbits during the period organogenesis caused adverse developmental outcomes including embryo-fetal mortality and malformations at maternal exposures (AUC) equivalent to or less than those in patients at the recommended doses. Advise pregnant women and females of reproductive potential of the potential risk to a fetus if SCSEMBLIX is used during pregnancy or if the patient becomes pregnant while taking SCSEMBLIX. Verify the pregnancy status of females of reproductive potential prior to starting treatment with SCSEMBLIX. Females of reproductive potential should use effective contraception during treatment with SCSEMBLIX and for 1 week after the last dose [see *Use in Specific Populations* (8.1, 8.3)].

6 ADVERSE REACTIONS

The following clinically significant adverse reactions can occur with SCSEMBLIX and are discussed in greater detail in other sections of the labeling:

- Myelosuppression [see *Warnings and Precautions* (5.1)]
- Pancreatic Toxicity [see *Warnings and Precautions* (5.2)]
- Hypertension [see *Warnings and Precautions* (5.3)]
- Hypersensitivity [see *Warnings and Precautions* (5.4)]
- Cardiovascular Toxicity [see *Warnings and Precautions* (5.5)]

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

The pooled safety population described in the WARNING AND PRECAUTIONS reflect exposure to SCSEMBLIX at 10 mg to 200 mg orally twice daily (between 0.25 to 5 times the recommended dosage for the 80 mg daily dosage and between 0.05 times and up to the recommended dosage for the 200 mg twice daily dosage) in 356 patients enrolled in one of two clinical trials, including patients with Ph+ CML in chronic (CP) receiving SCSEMBLIX as monotherapy: study CABL001A2301 (ASCEMBL) and study CABL001X2101 [see *Clinical Studies* (14)]. Among the 356 patients receiving SCSEMBLIX, the median duration of exposure to SCSEMBLIX was 89 weeks (range, 0.1 to 342 weeks).

Adverse Reactions in Patients with Ph+ CML-CP, Previously Treated with Two or More TKIs

The clinical trial randomized and treated 232 patients with Ph+ CML-CP, previously treated with two or more TKIs to receive SCEMBLIX 40 mg twice daily or bosutinib 500 mg once daily (ASCEMBL) [see *Clinical Studies (14.1)*]. The safety population (received at least 1 dose of SCEMBLIX) included 156 patients with Ph+ CML-CP, previously treated with two or more TKIs. Among patients who received SCEMBLIX, 83% were exposed for 24 weeks or longer and 67% were exposed for 48 weeks or longer.

Serious adverse reactions occurred in 15% of patients who received SCEMBLIX. Serious adverse reactions in $\geq 1\%$ included pyrexia (1.9%), cardiac failure congestive (1.3%), thrombocytopenia (1.3%), and urinary tract infection (1.3%). Two patients (1.3%) had a fatal adverse reaction, one each for mesenteric artery thrombosis and ischemic stroke.

Permanent discontinuation of SCEMBLIX due to an adverse reaction occurred in 7% of patients. Adverse reactions which resulted in permanent discontinuation of SCEMBLIX in $> 2\%$ of patients included thrombocytopenia (3.2%) and neutropenia (2.6%).

Dosage interruptions of SCEMBLIX due to an adverse reaction occurred in 38% of patients. Adverse reactions which required dosage interruption in $> 5\%$ of patients included thrombocytopenia (19%) and neutropenia (18%).

Dose reductions of SCEMBLIX due to an adverse reaction occurred in 7% of patients. Adverse reactions which required dose reductions in $> 1\%$ of patients included thrombocytopenia (4.5%) and neutropenia (1.3%).

The most common ($\geq 20\%$) adverse reactions in patients who received SCEMBLIX were upper respiratory tract infections and musculoskeletal pain.

The most common select laboratory abnormalities that worsened from baseline in $\geq 20\%$ of patients who received SCEMBLIX were platelet count decreased, triglycerides increased, neutrophil count decreased, hemoglobin decreased, creatine kinase increased, and alanine aminotransferase (ALT) increased.

Table 3 summarizes the adverse reactions in ASCEMBL.

Table 3: Adverse Reactions ($\geq 10\%$) in Patients with Ph+ CML in CP, Previously Treated with Two or More TKIs Who Received SCEMBLIX in ASCEMBL

Adverse Reaction	SCEMBLIX N = 156		Bosutinib N = 76	
	All Grades %	Grade 3 or 4 %	All Grades %	Grade 3 or 4 %
Infections and infestations				
Upper respiratory tract infection ^a	26	0.6	12	1.3
Musculoskeletal and connective tissue disorders				
Musculoskeletal pain ^b	22	2.6	16	1.3
Arthralgia	12	0	3.9	0
Nervous system disorders				
Headache	19	1.9	15	0
General disorders and administration-site conditions				
Fatigue ^c	17	0.6	11	1.3
Skin and subcutaneous tissue disorders				
Rash ^d	17	0.6	30	8
Vascular disorders				
Hypertension ^e	13	6	5	3.9
Gastrointestinal disorders				
Diarrhea ^f	12	0	71	11
Nausea	12	0.6	46	0
Abdominal pain ^g	10	0	24	2.6
Abbreviations: Ph+ CML in CP, Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic phase (CP); TKIs, tyrosine kinase inhibitors.				
^a Upper respiratory tract infection includes: nasopharyngitis, upper respiratory tract infection, rhinitis, pharyngitis, respiratory tract infection, and pharyngotonsillitis.				

^bMusculoskeletal pain includes: pain in extremity, back pain, myalgia, non-cardiac chest pain, neck pain, bone pain, spinal pain, arthritis, and musculoskeletal pain.

^cFatigue includes: fatigue and asthenia.

^dRash includes: rash, rash maculopapular, dermatitis acneiform, rash pustular, eczema, dermatitis, skin exfoliation, dermatitis exfoliative generalised, rash morbilliform, drug eruption, erythema multiforme, and rash erythematous.

^eHypertension includes: hypertension and hypertensive crisis.

^fDiarrhea includes: diarrhea and colitis.

^gAbdominal pain includes: abdominal pain, abdominal pain upper, abdominal discomfort, abdominal pain lower, abdominal tenderness, and epigastric discomfort.

Clinically relevant adverse reactions in < 10% of patients treated with SCEMBLIX in ASCSEMBL included: cough, dyspnea, pleural effusion, dizziness, neuropathy peripheral, edema, pyrexia, vomiting, constipation, dyslipidemia, decreased appetite, pruritus, urticaria, lower respiratory tract infection, influenza, urinary tract infection, pneumonia, hemorrhage, arrhythmia (including electrocardiogram QT prolonged), palpitations, cardiac failure congestive, vision blurred, dry eye, hypothyroidism, and febrile neutropenia.

Table 4 summarizes the laboratory abnormalities in ASCSEMBL.

Table 4: Select Laboratory Abnormalities (≥ 10%) That Worsened from Baseline in Patients with Ph+ CML in CP, Previously Treated with Two or More Tyrosine Kinase Inhibitors Who Received SCEMBLIX in ASCSEMBL

Laboratory Abnormality	SCEMBLIX ¹		Bosutinib ¹	
	All Grades %	Grade 3 or 4 %	All Grades %	Grade 3 or 4 %
Hematologic parameters				
Platelet count decreased	46	24	36	12
Neutrophil count decreased	39	17	33	13
Hemoglobin decreased	35	2	54	5
Lymphocyte count decreased	18	2	34	2.6
Biochemical parameters				
Triglycerides increased	44	5	29	2.6
Creatine kinase increased	27	2.6	22	5
Alanine aminotransferase (ALT) increased	23	0.6	50	16
Aspartate aminotransferase (AST) increased	19	1.9	46	7
Uric acid increased	19	6	17	2.6
Phosphate decreased	17	6	18	7
Lipase increased	14	3.9	18	7
Calcium corrected decreased	14	0.6	20	0
Creatinine increased	14	0	26	0
Amylase increased	12	1.3	13	0
Bilirubin increased	12	0	4.2	0
Cholesterol increased	11	0	8	0

Potassium decreased	10	0	9	0
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¹ The denominator used to calculate the rate for SCEMBLIX and bosutinib varied from 145 to 156 and 71 to 76, respectively, based on the number of patients with a baseline value and at least one post-treatment value.

CTCAE version 4.03.

Adverse Reactions in Patients with Ph+ CML-CP with the T315I Mutation

The single-arm clinical trial enrolled patients with Ph+ CML-CP with the T315I mutation [see *Clinical Studies (14.2)*]. The safety population (received at least 1 dose of SCEMBLIX) included 48 patients with Ph+ CML-CP with the T315I mutation who received 200 mg of SCEMBLIX twice daily. Among these patients, 83% were exposed for 24 weeks or longer and 75% were exposed for 48 weeks or longer.

Serious adverse reactions occurred in 23% of patients who received SCEMBLIX. Serious adverse reactions in > 1% included abdominal pain (4.2%), vomiting (4.2%), pneumonia (4.2%), musculoskeletal pain (2.1%), headache (2.1%), hemorrhage (2.1%), constipation (2.1%), arrhythmia (2.1%), and pleural effusion (2.1%).

Permanent discontinuation of SCEMBLIX due to an adverse reaction occurred in 10% of patients. Adverse reactions which resulted in permanent discontinuation of SCEMBLIX in > 2% of patients included pancreatic enzymes increased (2.1%).

Dosage interruptions of SCEMBLIX due to an adverse reaction occurred in 31% of patients. Adverse reactions which required dosage interruption in > 5% of patients included pancreatic enzymes increased (17%) and thrombocytopenia (8%).

Dose reductions of SCEMBLIX due to an adverse reaction occurred in 23% of patients. Adverse reactions which required dose reductions in > 1% of patients included pancreatic enzymes increased (10%), abdominal pain (4.2%), anemia (2.1%), blood bilirubin increased (2.1%), dizziness (2.1%), fatigue (2.1%), hepatic enzymes increased (2.1%), musculoskeletal pain (2.1%), nausea (2.1%), neutropenia (2.1%), pruritus (2.1%), and thrombocytopenia (2.1%).

The most common ($\geq 20\%$) adverse reactions in patients who received SCEMBLIX were musculoskeletal pain, fatigue, nausea, rash, and diarrhea.

The most common select laboratory abnormalities that worsened from baseline in $\geq 20\%$ of patients who received SCEMBLIX were alanine aminotransferase (ALT) increased, lipase increased, triglycerides increased, hemoglobin decreased, neutrophil count decreased, lymphocyte count decreased, phosphate decreased, aspartate aminotransferase (AST) increased, amylase increased, platelet count decreased, and bilirubin increased.

Table 5 summarizes adverse reactions in study X2101.

Table 5: Adverse Reactions ($\geq 10\%$) in Patients with Ph+ CML in CP with the T315I Mutation Who Received SCEMBLIX in X2101

Adverse Reaction	SCEMBLIX 200 mg twice daily N = 48	
	All Grades %	Grade 3 or 4 %
Musculoskeletal and connective tissue disorders		
Musculoskeletal pain ^a	42	4.2
Arthralgia	17	0
General disorders and administration-site conditions		
Fatigue ^b	31	2.1
Edema	10	4.2
Gastrointestinal disorders		
Nausea	27	0
Diarrhea	21	2.1
Vomiting	19	6
Abdominal pain ^c	17	8

Skin and subcutaneous tissue disorders		
Rash ^d	27	0
Pruritus	13	0
Nervous system disorders		
Headache ^e	19	2.1
Respiratory, thoracic, and mediastinal disorders		
Cough ^f	15	0
Vascular disorders		
Hemorrhage ^g	15	2.1
Hypertension ^h	13	8
Infections and infestations		
Upper respiratory tract infection ⁱ	13	0

^aMusculoskeletal pain includes: pain in extremity, back pain, myalgia, musculoskeletal pain, non-cardiac chest pain, bone pain, arthritis, and musculoskeletal chest pain.
^bFatigue includes: fatigue and asthenia.
^cAbdominal pain includes: abdominal pain and hepatic pain.
^dRash includes: rash, rash maculopapular, dermatitis acneiform, eczema, rash papular, skin exfoliation, and dyshidrotic eczema.
^eHeadache includes: headache and migraine.
^fCough includes: cough and productive cough.
^gHemorrhage includes: epistaxis, ear hemorrhage, mouth hemorrhage, post procedural hemorrhage, skin hemorrhage, and vaginal hemorrhage.
^hHypertension includes: hypertension and hypertensive crisis.
ⁱUpper respiratory tract infection includes: upper respiratory tract infection, nasopharyngitis, rhinitis, and pharyngitis.

Clinically relevant adverse reactions in < 10% of patients treated with SCEMBLIX in X2101 included: constipation, pancreatitis, pyrexia, dizziness, neuropathy peripheral, pneumonia, lower respiratory tract infection, dyspnea, pleural effusion, dry eye, vision blurred, arrhythmia, palpitations, cardiac failure congestive, decreased appetite, dyslipidemia, and urticaria.

Table 6 summarizes laboratory abnormalities in X2101.

Table 6: Select Laboratory Abnormalities (≥ 10%) That Worsened from Baseline in Patients with Ph+ CML in CP with the T315I Mutation in X2101

Laboratory Abnormality	SCEMBLIX¹ 200 mg twice daily	
	All Grades %	Grade 3-4 %
Hematologic parameters		
Hemoglobin decreased	44	4.2
Neutrophil count decreased	44	15
Lymphocyte count decreased	42	4.2
Platelet count decreased	25	15
Biochemical parameters		
Alanine aminotransferase (ALT) increased	48	6
Potassium increased	48	2.1
Triglycerides increased	46	2.1
Lipase increased	46	21
Phosphate decreased	40	6
Uric acid increased	40	4.2

Aspartate aminotransferase (AST) increased	35	2.1
Calcium corrected decreased	33	0
Creatinine increased	31	0
Amylase increased	29	10
Bilirubin increased	23	0
Cholesterol increased	15	0
Alkaline phosphatase (ALP) increased	13	0

[†]The denominator used to calculate the rate was 48 based on the number of patients with a baseline value and a least one post-treatment value.

CTCAE version 4.03.

7 DRUG INTERACTIONS

7.1 Effect of Other Drugs on SCEMBLIX

Strong CYP3A4 Inhibitors

Asciminib is a CYP3A4 substrate. Concomitant use of SCEMBLIX with a strong CYP3A4 inhibitor increases both the asciminib C_{max} and AUC, which may increase the risk of adverse reactions [see *Clinical Pharmacology (12.3)*]. Closely monitor for adverse reactions in patients treated with SCEMBLIX at 200 mg twice daily with concomitant use of strong CYP3A4 inhibitors.

Itraconazole Oral Solution Containing Hydroxypropyl- β -cyclodextrin

Concomitant use of SCEMBLIX with itraconazole oral solution containing hydroxypropyl- β -cyclodextrin decreases asciminib C_{max} and AUC, which may reduce SCEMBLIX efficacy [see *Clinical Pharmacology (12.3)*]. Avoid coadministration of SCEMBLIX at all recommended doses with itraconazole oral solution containing hydroxypropyl- β -cyclodextrin.

7.2 Effect of SCEMBLIX on Other Drugs

Certain CYP3A4 Substrates

Asciminib is a CYP3A4 inhibitor. Concomitant use of SCEMBLIX increases the C_{max} and AUC of CYP3A4 substrates, which may increase the risk of adverse reactions of these substrates [see *Clinical Pharmacology (12.3)*].

Closely monitor for adverse reactions in patients treated with SCEMBLIX at 80 mg total daily dose with concomitant use of certain CYP3A4 substrates, where minimal concentration changes may lead to serious adverse reactions. Avoid coadministration of SCEMBLIX at 200 mg twice daily with certain CYP3A4 substrates, where minimal concentration changes may lead to serious adverse reactions.

CYP2C9 Substrates

Asciminib is a CYP2C9 inhibitor. Concomitant use of SCEMBLIX increases the C_{max} and AUC of CYP2C9 substrates, which may increase the risk of adverse reactions of these substrates [see *Clinical Pharmacology (12.3)*].

Avoid coadministration of SCEMBLIX at 80 mg total daily dose with certain CYP2C9 substrates, where minimal concentration changes may lead to serious adverse reactions. If coadministration is unavoidable, reduce the CYP2C9 substrate dosage as recommended in its prescribing information.

Avoid coadministration of SCEMBLIX at 200 mg twice daily with sensitive CYP2C9 substrates and certain CYP2C9 substrates, where minimal concentration changes may lead to serious adverse reactions. If coadministration is unavoidable, consider alternative therapy with non-CYP2C9 substrate.

Certain P-gp Substrates

Asciminib is a P-gp inhibitor. Concomitant use of SCEMBLIX increases the plasma concentrations of P-gp substrates, which may increase the risk of adverse reactions of these substrates [see *Clinical Pharmacology (12.3)*].

Closely monitor for adverse reactions in patients treated with SCEMBLIX at all recommended doses with concomitant use of P-gp substrates, where minimal concentration changes may lead to serious toxicities.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Based on findings from animal studies and the mechanism of action, SCEMBLIX can cause embryo-fetal harm when administered to a pregnant woman [see *Clinical Pharmacology (12.1)*]. There are no available data on SCEMBLIX use in pregnant women to evaluate a drug associated risk.

Animal reproduction studies in pregnant rats and rabbits demonstrated that oral administration of asciminib during organogenesis induced structural abnormalities, embryo-fetal mortality, and alterations to growth (see *Data*).

Advise pregnant women and females of reproductive potential of the potential risk to a fetus.

In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

Data

Animal Data

In embryo-fetal development studies, pregnant animals received oral doses of asciminib at 25, 150, and 600 mg/kg/day in rats and at 15, 50, and 300 mg/kg/day in rabbits during the period of organogenesis.

In rats, maternal toxicity at the asciminib dose of 600 mg/kg/day resulted in the early termination of the dose group; a complete embryo-fetal examination was not conducted for this group. Adverse embryo-fetal findings were observed at 25 and 150 mg/kg; these doses did not cause maternal toxicities. Increases in fetal weights at 25 and 150 mg/kg/day were observed, which may be related to increased ossification (i.e. increased rate of development). Malformations were evident at 150 mg/kg and included cleft palate, anasarca (edema), and cardiac abnormalities. Additional fetal findings included urinary tract and skeletal variations, observed primarily at 150 mg/kg/day. At the of 25 mg/kg/day, the area under the curve (AUC) exposures were equivalent to or below those achieved in patients at the 40 mg twice daily or 80 mg once daily doses, respectively. At the dose of 25 mg/kg/day, the AUC exposures were below those achieved in patients at the 200 mg twice daily dose.

In rabbits, maternal toxicities at the asciminib dose of 300 mg/kg/day resulted in the early termination of the dose group; a complete embryo-fetal examination was not conducted for this group. Adverse embryo-fetal findings were observed at 50 mg/kg; this dose did not cause maternal toxicities. Findings at the 50 mg/kg dose included increases in early resorptions and post-implantation loss, decreases in the number of live fetuses, and cardiac malformations. At the dose of 50 mg/kg/day, the AUC exposures were 4-fold those achieved in patients at the 40 mg twice daily or 80 mg once daily doses. At the dose of 50 mg/kg/day, the AUC exposures were below those achieved in patients at the 200 mg twice daily dose.

8.2 Lactation

Risk Summary

There are no data on the presence of asciminib or its metabolites in human milk, the effects on the breastfed child, or milk production.

Because of the potential for serious adverse reactions in the breastfed child, advise women not to breastfeed during treatment with SCEMBLIX and for 1 week after the last dose.

8.3 Females and Males of Reproductive Potential

Based on findings from animal studies, SCEMBLIX can cause embryo-fetal harm when administered to a pregnant woman [see *Use in Specific Populations (8.1)*].

Pregnancy Testing

Verify the pregnancy status of females of reproductive potential prior to starting treatment with SCEMBLIX.

Contraception

Females

Females of reproductive potential should use effective contraception during treatment with SCEMBLIX and for 1 week after the last dose.

Infertility

Based on findings in animals, SCEMBLIX may impair fertility in females of reproductive potential [see *Nonclinical Toxicology (13.1)*]. The reversibility of the effect on fertility is unknown

8.4 Pediatric Use

The safety and efficacy of SCEMBLIX in pediatric patients have not been established.

8.5 Geriatric Use

In the ASCEMBL study, 44 of the 233 (19%) patients were 65 years of age or older and 6 (2.6%) were 75 years of age or older.

In the X2101 study, 16 of the 48 (33%) patients with the T315I mutation were 65 years of age or older and 4 (8%) were 75 years of age or older.

Overall, no differences in safety or efficacy of SCEMBLIX were observed between patients 65 years of age or older compared to younger patients. There is an insufficient number of patients 75 years of age or older to assess whether there are differences in safety or efficacy.

8.6 Renal Impairment

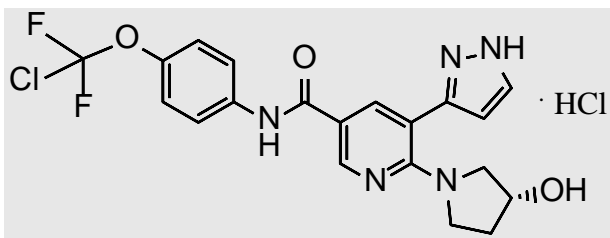
No dose adjustment is required for patients with mild to severe renal impairment (estimated glomerular filtration rate [eGFR] 15 to 89 mL/min/1.73 m²) and not requiring dialysis receiving SCEMBLIX [see *Clinical Pharmacology (12.3)*].

8.7 Hepatic Impairment

No dose adjustment is required for patients with mild [total bilirubin \leq upper limit of normal (ULN) and aspartate aminotransferase (AST) $>$ ULN or total bilirubin $>$ 1 to 1.5 times ULN and any AST] to severe hepatic impairment (total bilirubin $>$ 3 times ULN and any AST) receiving SCEMBLIX [see *Clinical Pharmacology (12.3)*].

11 DESCRIPTION

SCEMBLIX (asciminib) is a kinase inhibitor. The chemical name of the drug substance is *N*-[4-(Chlorodifluoromethoxy)phenyl]-6-[(3*R*)-3-hydroxypyrrolidin-1-yl]-5-(1*H*-pyrazol-3-yl)pyridine-3-carboxamide hydrochloride (1/1). Asciminib hydrochloride is a white to slightly yellow powder. The molecular formula of asciminib hydrochloride is C₂₀H₁₈ClF₂N₅O₃·HCl, and the relative molecular mass is 486.30 g/mol for the hydrochloride salt and 449.84 g/mol for the free base. The chemical structure of asciminib hydrochloride is shown below:



SCEMBLIX film-coated tablets are supplied for oral use with two strengths that contain 20 mg and 40 mg of asciminib (equivalent to 21.62 mg and 43.24 mg, respectively, of asciminib hydrochloride). The tablets contain colloidal silicon dioxide, croscarmellose sodium, ferric oxide, hydroxypropyl cellulose, lactose monohydrate, lecithin, magnesium stearate, microcrystalline cellulose, polyvinyl alcohol, talc, titanium dioxide, and xanthan gum. The 20 mg tablets contain ferric oxide, yellow and ferric oxide, red. The 40 mg tablets contain ferrousferrous oxide and ferric oxide, red.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Asciminib is an ABL/BCR-ABL1 tyrosine kinase inhibitor. Asciminib inhibits the ABL1 kinase activity of the BCR-ABL1 fusion protein, by binding to the ABL myristoyl pocket. In studies conducted in vitro or in animal models of CML, asciminib showed activity against wild-type BCR-ABL1 and several mutant forms of the kinase, including the T315I mutation.

12.2 Pharmacodynamics

Exposure-Response Relationships

Over asciminib dosages of 10 mg to 200 mg twice daily (0.25 to 5 times the recommended 80 mg daily dosage), a lower exposure was associated with a smaller decrease in *BCR-ABL1* level and a lower MMR rate at Week 24.

Over asciminib dosages of 10 mg to 280 mg twice daily (0.25 to 7 times the recommended 80 mg daily dosage), a higher exposure was associated with slightly higher incidence of some adverse reactions (e.g., Grade ≥ 3 lipase increase, Grade ≥ 3 hemoglobin decrease, Grade ≥ 2 ALT increase, Grade ≥ 2 AST increase, Grade ≥ 2 bilirubin increase, and any grade lipase increase).

Cardiac Electrophysiology

Asciminib does not cause a large mean increase in QTc interval (i.e., >20 msec) at the maximum recommended clinical dosage (200 mg twice daily). Based on available clinical data, small mean QTc increase (<10 msec) cannot be excluded.

12.3 Pharmacokinetics

Asciminib steady-state exposure (AUC and C_{\max}) increase slightly more than dose proportional across the dose range of 10 to 200 mg (0.25 to 5 times the recommended 80 mg daily dosage) administered once or twice daily.

Pharmacokinetic parameters are presented as geometric mean (CV%) unless otherwise stated. The steady state C_{\max} and AUC_{τ} of asciminib at recommended dosages are listed in Table 7.

Table 7: Steady State^a Asciminib Exposure at Recommended Dosages

Asciminib Dosage	C_{\max} (ng/mL)	AUC_{τ} ^b (ng*h/mL)	Accumulation Ratio
80 mg once daily	1781 (23%)	15112 (28%)	1.30
40 mg twice daily	793 (49%)	5262 (48%)	1.65
200 mg twice daily	5642 (40%)	37547 (41%)	1.92

^aSteady state is achieved within 3 days.

^b AUC_{τ} represents AUC_{0-12h} for twice daily dosing and AUC_{0-24h} for once daily dosing.

Absorption

The median (range) T_{\max} of asciminib is 2.5 hours (2 to 3 hours).

Effect of Food

The AUC and C_{\max} of asciminib decreased by 62% and 68%, respectively, with a high-fat meal (1000 calories, 50% fat) and by 30% and 35%, respectively, with a low-fat meal (400 calories, 25% fat) compared to the fasted state following administration of SCEMBLIX.

Distribution

The apparent volume of distribution of asciminib at steady state is 151 L (135%). Asciminib is the main circulating component in plasma (93% of the administered dose).

Asciminib is 97% bound to human plasma proteins *in vitro*.

Elimination

The total apparent clearance of asciminib is 6.7 L/hour (48%) at 40 mg twice daily and 80 mg once daily, and 4.1 L/hour (38%) at 200 mg twice daily. The terminal elimination half-life of asciminib is 5.5 hours (38%) at 40 mg twice daily and 80 mg once daily, and 9.0 hours (33%) at 200 mg twice daily.

Metabolism

Asciminib is metabolized by CYP3A4-mediated oxidation, UGT2B7- and UGT2B17-mediated glucuronidation.

Excretion

Eighty percent (57% as unchanged) and 11% (2.5% as unchanged) of the asciminib dose were recovered in the feces and in the urine of healthy subjects, respectively, following oral administration of a single 80 mg dose of radio-labeled asciminib.

Asciminib is eliminated by biliary secretion via breast cancer-resistant protein (BCRP).

Specific Populations

No clinically significant differences in the pharmacokinetics of asciminib were observed based on sex, age (20 to 88 years), race (Asian 20%, White 70%, Black/African American 4%), or body weight (42 - 184 kg), mild to moderate renal impairment (eGFR 30 to 89 mL/min/1.73 m²), or mild (total bilirubin ≤ ULN and AST > ULN or total bilirubin > 1 to 1.5 times ULN and any AST) to moderate (total bilirubin > 1.5 to 3 times ULN and any AST) hepatic impairment.

Patients with Renal Impairment

Asciminib AUC_{inf} and C_{max} are increased by 57% and 6%, respectively, in subjects with eGFR between 13 to < 30 mL/min/1.73 m² and not requiring dialysis compared to subjects with normal renal function (eGFR ≥ 90 mL/min/1.73 m²) following oral administration of a single 40 mg dose of SCEMBLIX. The exposure changes in patients with severe renal impairment are not considered clinically meaningful.

Patients with Hepatic Impairment

Asciminib AUC_{inf} and C_{max} are increased by 33% and 4%, respectively, in subjects with severe hepatic impairment (total bilirubin > 3 times ULN and any AST), compared to subjects with normal hepatic function (total bilirubin ≤ ULN and AST ≤ ULN) following oral administration of a single 40 mg dose of SCEMBLIX. The exposure changes are not considered clinically meaningful.

Drug Interaction Studies

Clinical Studies and Model-Informed Approaches

Drugs That Affect Asciminib Plasma Concentration

Strong CYP3A Inhibitors: The asciminib AUC_{inf} and C_{max} increased by 36% and 19%, respectively, following coadministration of a single SCEMBLIX dose of 40 mg with a strong CYP3A4 inhibitor (clarithromycin). No clinically significant differences in the pharmacokinetics of asciminib were observed when coadministered with itraconazole, which is also a strong CYP3A4 inhibitor.

Strong CYP3A Inducers: Concomitant use of strong CYP3A inducers with SCEMBLIX has not been fully characterized.

Itraconazole Oral Solution: Coadministration of multiple doses of itraconazole oral solution containing hydroxypropyl-β-cyclodextrin with a single SCEMBLIX dose of 40 mg decreased asciminib AUC_{inf} and C_{max} by 40% and 50%, respectively. Concomitant use of oral products containing hydroxypropyl-β-cyclodextrin with SCEMBLIX other than itraconazole oral solution has not been fully characterized.

Imatinib: The asciminib AUC_{inf} and C_{max} increase by 108% and 59%, respectively following coadministration of a single SCEMBLIX dose of 40 mg with imatinib (an inhibitor of BCRP, CYP3A4, UGT2B17 and UGT1A3/4). The exposure changes are not considered clinically meaningful. Concomitant use of imatinib with SCEMBLIX at 200 mg twice daily has not been fully characterized.

Other Drugs: No clinically significant differences in the pharmacokinetics of asciminib were observed when coadministered with rabeprazole (acid-reducing agent) and quinidine (P-gp inhibitor).

Drugs That are Affected by Asciminib

CYP3A4 Substrates: The midazolam AUC_{inf} and C_{max} increased by 28% and 11%, respectively, following coadministration of a CYP3A4 substrate (midazolam) with SCEMBLIX 40 mg twice daily. The midazolam AUC_{inf} and C_{max} increased by 24% and 17%, respectively, following coadministration with SCEMBLIX at 80 mg once daily and 88% and 58%, respectively, at 200 mg twice daily.

CYP2C9 Substrates: The S-warfarin AUC_{inf} and C_{max} increased by 41% and 8%, respectively, following coadministration of CYP2C9 substrate (warfarin) with SCEMBLIX at 40 mg twice daily. The S-warfarin AUC_{inf} and C_{max} increased by 52% and 4%, respectively, following coadministration with SCEMBLIX at 80 mg once daily and 314% and 7%, respectively, at 200 mg twice daily.

CYP2C8 Substrates: The repaglinide (substrate of CYP2C8, CYP3A4, and OATP1B) AUC_{inf} and C_{max} increased by 8% and 14%, respectively, following coadministration of repaglinide with SCEMBLIX 40 mg twice daily. The repaglinide AUC_{inf} and C_{max} increased by 12% and 8%, respectively, following coadministration with SCEMBLIX at 80 mg once daily and 42% and 25%, respectively, at 200 mg twice daily. The rosiglitazone (substrate of CYP2C8 and CYP2C9) AUC_{inf} and C_{max} increased by 20% and 3%, respectively, following coadministration of rosiglitazone with SCEMBLIX 40

mg twice daily. The rosiglitazone AUC_{inf} and C_{max} increased by 24% and 2%, respectively, following coadministration with SCEMBLIX at 80 mg once daily and 66% and 8%, respectively, at 200 mg twice daily.

P-gp Substrates: Coadministration of SCEMBLIX with a drug that is a substrate of P-gp may result in a clinically relevant increase in the plasma concentrations of P-gp substrates, where minimal concentration changes may lead to serious toxicities.

In Vitro Studies

CYP450 and UGT Enzymes

Asciminib may reversibly inhibit UGT1A1 at plasma concentrations reached at a total daily dose of 80 mg and 200 mg twice daily. In addition, asciminib may reversibly inhibit CYP2C19 at concentrations reached at 200 mg twice daily dose.

Transporter Systems

Asciminib is a substrate of BCRP and P-gp. Asciminib inhibits BCRP, P-gp, OATP1B1, OATP1B3, and OCT1.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Carcinogenicity studies have not been conducted with asciminib.

Asciminib was not genotoxic in an *in vitro* bacterial mutagenicity (Ames) assay, an *in vitro* micronucleus assay in human peripheral blood lymphocytes (HPBL) or an *in vivo* rat peripheral blood reticulocyte micronucleus assay.

In a combined male and female fertility and early embryonic development study in rats, animals were administered asciminib doses of 10, 50, or 200 mg/kg/day orally. Male animals were dosed once daily for at least 28 days prior to mating, during the 2-week mating period, and until terminal necropsy (Days 63-67). Female animals were dosed once daily for the 2-week pre-mating period, during the 2-week mating period, and through gestation day (GD) 6. Decreased sperm count and motility were observed at 200 mg/kg/day. While there were no effects on fertility indices or conception rates, a decreased mean number of live embryos was observed at 200 mg/kg/day and was attributed to a lower number of implantations and an increased number of early resorptions. Increased early resorptions were also observed in the embryo-fetal development study in rabbits [see *Use in Specific Populations (8.1)*].

At the dose of 200 mg/kg, the AUC exposures were approximately 19-fold, 13-fold, or 2-fold higher than those achieved in patients at the 40 mg twice daily, 80 mg once daily, or 200 mg twice daily doses, respectively.

14 CLINICAL STUDIES

14.1 Ph+ CML-CP, Previously Treated With Two or More TKIs

The efficacy of SCEMBLIX in the treatment of patients with Ph+ CML in chronic phase (Ph+ CML-CP), previously treated with two or more tyrosine kinase inhibitors was evaluated in the multi-center, randomized, active-controlled, and open-label study ASCEMBL (NCT 03106779).

In this study, a total of 233 patients were randomized in a 2:1 ratio and stratified according to major cytogenetic response (MCyR) status to receive either SCEMBLIX 40 mg twice daily (N = 157) or bosutinib 500 mg once daily (N = 76). Patients continued treatment until unacceptable toxicity or treatment failure occurred.

Patients were 52% female and 48% male with a median age of 52 years (range, 19 to 83 years). Of the 233 patients, 19% were 65 years or older, while 2.6% were 75 years or older. Patients were White (75%), Asian (14%), and Black or African American (4.3%). Of the 233 patients, 81% and 18% had Eastern Cooperative Oncology Group (ECOG) performance status 0 or 1, respectively. Patients who had previously received 2, 3, 4, or 5 or more prior lines of TKIs were 48%, 31%, 15%, and 6%, respectively. The median duration of treatment was 67 weeks (range, 0.1 to 162 weeks) for patients receiving SCEMBLIX and 30 weeks (range, 1 to 149 weeks) for patients receiving bosutinib.

The main efficacy outcomes from ASCEMBL are summarized in Table 8.

Table 8: Efficacy Results in Patients With Ph+ CML-CP, Previously Treated With Two or More Tyrosine Kinase Inhibitors (ASCEMBL)

	SCEMBLIX 40 mg twice daily	Bosutinib 500 mg once daily	Difference (95% CI)	p-value
MMR rate, % (95% CI) at 24 weeks	N = 157 25 (19, 33)	N = 76 13 (6.5, 23)	12 ^a (2.2, 22)	0.029 ^b
CCyR rate, % (95% CI) at 24 weeks	N = 103 ^c 41 (31, 51)	N = 62 ^c 24 (14, 37)	17 (3.6, 31)	

Abbreviations: MMR, major molecular response ($BCR-ABL1^{IS} \leq 0.1\%$); CCyR, complete cytogenetic response (0% of Philadelphia-positive metaphases in bone marrow aspirate with at least 20 examined).
^aEstimated using a common risk difference stratified by baseline major cytogenetic response status.
^bEstimated using a Cochrane-Mantel-Haenszel two-sided test stratified by baseline major cytogenetic response status.
^cCCyR analysis based on patients who were not in CCyR at baseline.

The MMR rate at 48 weeks was 29% (95% CI: 22, 37) in patients receiving SCEMBLIX and 13% (95% CI: 6.5, 23) in patients receiving bosutinib. With a median duration of follow-up of 20 months (range: 1 day to 36 months), the median duration of response had not yet been reached for patients with MMR at any time.

14.2 Ph+ CML-CP with the T315I mutation

The efficacy of SCEMBLIX in the treatment of patients with Ph+ CML-CP with the T315I mutation was evaluated in a multi-center open-label study CABL001X2101 (NCT02081378). Testing for T315I mutation utilized a qualitative p210 BCR-ABL mutation test using Sanger Sequencing.

Efficacy was based on 45 patients with Ph+ CML-CP with the T315I mutation who received SCEMBLIX at a dose of 200 mg twice daily. Patients continued treatment until unacceptable toxicity or treatment failure occurred.

Of the 45 patients, 80% were male and 20% female; 31% were 65 years or older, while 9% were 75 years or older with a median age of 54 years (range, 26 to 86 years). The patients were White (47%), Asian (27%), and Black or African American (2.2%), and 24% were unreported or unknown. Seventy-three percent and 27% of patients had ECOG performance status 0 and 1, respectively. Patients who had previously received 1, 2, 3, 4, and 5 or more TKIs were 18%, 31%, 36%, 13%, and 2.2%, respectively.

MMR was achieved by 24 weeks in 42% (19/45, 95% CI: 28% to 58%) of the 45 patients treated with SCEMBLIX. MMR was achieved by 96 weeks in 49% (22/45, 95% CI: 34% to 64%) of the 45 patients treated with SCEMBLIX. The median duration of treatment was 108 weeks (range, 2 to 215 weeks).

16 HOW SUPPLIED/STORAGE AND HANDLING

SCEMBLIX tablets are available as:

Table 9: SCEMBLIX Package Configurations and NDC Numbers

Package Configuration	Tablet Strength	NDC Number
Bottle of 60 tablets	20 mg	NDC 0078-1091-20
Bottle of 60 tablets	40 mg	NDC 0078-1098-20
Carton containing 5 bottles. Each bottle contains 60 tablets.	40 mg	NDC 0078-1098-30

- SCEMBLIX (asciminib) 20 mg film-coated tablets are supplied as pale yellow, unscored, round, biconvex, with beveled edges, film-coated tablet containing 20 mg of asciminib (equivalent to 21.62 mg of asciminib HCl). Each tablet is debossed with “20” on one side and “Novartis logo” on the other side.

- SCEMBLIX (asciminib) 40 mg film-coated tablets are supplied as violet white, unscored, round, biconvex, with beveled edges, film-coated tablet containing 40 mg of asciminib (equivalent to 43.24 mg of asciminib HCl). Each tablet is debossed with “40” on one side and “Novartis logo” on the other side.

Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C and 30°C (59°F and 86°F) [see USP Controlled Room Temperature]. Dispense and store in the original container in order to protect from moisture.

17 PATIENT COUNSELING INFORMATION

Advise the patient to read the FDA-approved patient labeling (Patient Information).

Myelosuppression

Inform patients of the possibility of developing low blood cell counts. Advise patients to immediately report fever, any suggestion of infection, or signs or symptoms suggestive of bleeding or easy bruising [see *Warnings and Precautions (5.1)*].

Pancreatic Toxicity

Inform patients of the possibility of developing pancreatitis that may be accompanied by nausea, vomiting, severe abdominal pain, or abdominal discomfort, and to promptly report these symptoms [see *Warnings and Precautions (5.2)*].

Hypertension

Inform patients of the possibility of developing hypertension. Advise patients to contact their healthcare provider for elevated blood pressure or if symptoms of hypertension occur including confusion, headache, dizziness, chest pain, or shortness of breath [see *Warnings and Precautions (5.3)*].

Hypersensitivity

Advise the patient to discontinue SCEMBLIX and seek immediate medical attention if any signs or symptoms of a hypersensitivity reaction, such as rash, edema, or bronchospasm occur [see *Warnings and Precautions (5.4)*].

Cardiovascular Toxicity

Inform patients of the possibility of the occurrence of cardiovascular toxicity, especially those with a history of cardiovascular risk factors. Advise patients to immediately contact their healthcare provider or get medical help if they develop cardiovascular signs and symptoms [see *Warnings and Precautions (5.5)*].

Embryo-Fetal Toxicity

Advise females to inform their healthcare provider if they are pregnant or become pregnant. Inform female patients of the potential risk to a fetus [see *Use in Specific Populations (8.1)*].

Advise females of reproductive potential to use effective contraception during treatment and for 1 week after receiving the last dose of SCEMBLIX [see *Warnings and Precautions (5.6)*, *Use in Specific Populations (8.1, 8.3)*].

Lactation

Advise women not to breastfeed during treatment with SCEMBLIX and for 1 week after the last dose [see *Use in Specific Populations (8.2)*].

Drug Interactions

Advise patients that SCEMBLIX and certain other medicines, including over the counter medications or herbal supplements, can interact with each other and may alter the effects of SCEMBLIX [see *Drug Interactions (7)*].

Instructions for Taking SCEMBLIX

Advise patients to take SCEMBLIX exactly as prescribed and not to change their dose or schedule or to stop taking SCEMBLIX unless they are told to do so by their healthcare provider.

Advise patients to take SCEMBLIX orally without food. Advise patients to avoid food for at least 2 hours before and 1 hour after taking SCEMBLIX. SCEMBLIX tablets should be swallowed whole. Patients should not break, crush, or chew the tablets [see *Dosage and Administration (2.5)*].

Advise patients that if they take SCEMBLIX once daily and miss a dose by more than 12 hours to skip the missed dose.
Advise patients that if they take SCEMBLIX twice daily and miss a dose by more than 6 hours to skip the missed dose.
Advise patients to take the next dose as scheduled [*see Dosage and Administration (2.3)*].

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PATIENT INFORMATION

SCEMBLIX® (sem blix) (asciminib) tablets

What is SCEMBLIX?

SCEMBLIX is a prescription medicine used to treat adults with:

- Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic phase (CP), previously treated with 2 or more tyrosine kinase inhibitor (TKI) medicines.
- Ph+ CML in CP with the T315I mutation.

It is not known if SCEMBLIX is safe and effective in children.

Before taking SCEMBLIX, tell your healthcare provider about all of your medical conditions, including if you:

- have a history of inflammation of your pancreas (pancreatitis)
- have a history of heart problems or blood clots in your arteries and veins (types of blood vessels)
- are pregnant or plan to become pregnant. SCEMBLIX can harm your unborn baby.
 - Your healthcare provider will do a pregnancy test before you start treatment with SCEMBLIX.
 - **Females** who are able to become pregnant should use effective birth control during treatment and for 1 week after your last dose of SCEMBLIX. Talk to your healthcare provider about birth control methods that may be right for you.
 - Tell your healthcare provider right away if you become pregnant or think you may be pregnant during treatment with SCEMBLIX.
- are breastfeeding or plan to breastfeed. It is not known if SCEMBLIX passes into your breast milk. Do not breastfeed during treatment and for 1 week after your last dose of SCEMBLIX.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. SCEMBLIX and other medicines may affect each other causing side effects.

How should I take SCEMBLIX?

- Take SCEMBLIX exactly as your healthcare provider tells you.
- **Do not** change your dose or schedule or stop taking SCEMBLIX unless your healthcare provider tells you to.
- Take SCEMBLIX without food. **You should avoid eating** for at least 2 hours before and 1 hour after taking SCEMBLIX.
- Swallow SCEMBLIX tablets whole. **Do not** break, crush, or chew SCEMBLIX tablets.
- If you take SCEMBLIX **1 time a day** and miss a dose by more than 12 hours, skip the missed dose and take your next dose at your regular time.
- If you take SCEMBLIX **2 times a day** and miss a dose by more than 6 hours, skip the missed dose and take your next dose at your regular time.

What are the possible side effects of SCEMBLIX?

SCEMBLIX may cause serious side effects, including:

- **Low blood cell counts.** SCEMBLIX may cause low platelet counts (thrombocytopenia), low white blood cell counts (neutropenia), and low red blood cell counts (anemia). Your healthcare provider will do blood tests to check your blood cell counts every 2 weeks for the first 3 months of treatment and then monthly or as needed during treatment with SCEMBLIX. Tell your healthcare provider right away if you have unexpected bleeding or easy bruising, blood in your urine or stools, fever, or any signs of an infection.
- **Pancreas problems.** SCEMBLIX may increase enzymes in your blood called amylase and lipase, which may be a sign of pancreatitis. Your healthcare provider may do blood tests monthly or as needed during treatment with SCEMBLIX to check for problems with your pancreas. Tell your healthcare provider right away if you have sudden stomach-area pain or discomfort, nausea, or vomiting.
- **High blood pressure.** Your healthcare provider may check your blood pressure and treat any high blood pressure during treatment with SCEMBLIX as needed. Tell your healthcare provider if you develop elevated blood pressure or symptoms of high blood pressure including confusion, headaches, dizziness, chest pain, or shortness of breath.
- **Allergic reaction.** Stop taking SCEMBLIX and get medical help right away if you get any signs or symptoms of an allergic reaction, including:
 - trouble breathing or swallowing
 - swelling of the face, lips, or tongue
 - skin rash or flushing of your skin
 - feeling dizzy or faint
 - fever
 - fast heartbeat

- **Heart and blood vessel (cardiovascular) problems.** SCEMBLIX may cause heart and blood vessel problems, including heart attack, stroke, blood clots or blockage of your arteries, heart failure, and abnormal heartbeat, which can be serious and may sometimes lead to death. These heart and blood vessel problems can happen in people with risk factors or a history of these problems, and/or previously treated with other TKI medicines. Your healthcare provider may monitor you for heart and blood vessel problems and treat you as needed during treatment with SCEMBLIX. Tell your healthcare provider or get medical help right away if you get:

- shortness of breath
- chest pain or pressure
- feeling like your heart is beating too fast or you feel abnormal heartbeats
- swelling in your ankles or feet
- dizziness
- weight gain
- numbness or weakness on one side of your body
- decreased vision or loss of vision
- trouble talking
- pain in your arms, legs, back, neck or jaw
- headache
- severe stomach area pain

The most common side effects of SCEMBLIX include:

- nose, throat, or sinus (upper respiratory tract) infections
- muscle, bone, or joint pain
- tiredness
- nausea
- rash
- diarrhea
- decreased blood platelet counts, white blood cell counts, and red blood cell counts
- increased blood fat (triglycerides) levels
- increased blood creatine kinase levels
- increased blood liver enzyme levels
- increased blood pancreas enzyme (amylase and lipase) levels

Your healthcare provider may change your dose or temporarily or permanently stop treatment with SCEMBLIX if you have certain side effects.

SCEMBLIX may cause fertility problems in females. This may affect your ability to have a child. Talk to your healthcare provider if this is a concern for you.

These are not all of the possible side effects of SCEMBLIX.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

How should I store SCEMBLIX?

- Store SCEMBLIX at room temperature between 68°F to 77°F (20°C to 25°C).
- Store SCEMBLIX in the original container to protect it from moisture.

Keep SCEMBLIX and all medicines out of the reach of children.

General information about the safe and effective use of SCEMBLIX.

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use SCEMBLIX for a condition for which it was not prescribed. Do not give SCEMBLIX to other people, even if they have the same symptoms you have. It may harm them. You can ask your pharmacist or healthcare provider for more information about SCEMBLIX that is written for health professionals.

What are the ingredients in SCEMBLIX?

Active ingredient: asciminib

Inactive ingredients: colloidal silicon dioxide, croscarmellose sodium, ferric oxide, hydroxypropyl cellulose, lactose monohydrate, lecithin, magnesium stearate, microcrystalline cellulose, polyvinyl alcohol, talc, titanium dioxide, and xanthan gum. The 20 mg tablets contain ferric oxide, yellow and ferric oxide, red. The 40 mg tablets contain ferrousferrous oxide and ferric oxide, red.

Distributed by: Novartis Pharmaceuticals Corporation, One Health Plaza, East Hanover, New Jersey 07936
For more information, go to www.SCEMBLIX.com or call 1-888-669-6682.

This Patient Information has been approved by the U.S. Food and Drug Administration.

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