

PEPAXTO- melphalan flufenamide injection, powder, lyophilized, for solution Oncopeptides, AB

HIGHLIGHTS OF PRESCRIBING INFORMATION

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

PEPAXTO® (melphalan flufenamide) for injection, for intravenous use
Initial U.S. Approval: 2021

INDICATIONS AND USAGE

PEPAXTO is an alkylating drug indicated in combination with dexamethasone, for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy and whose disease is refractory to at least one proteasome inhibitor, one immunomodulatory agent, and one CD38-directed monoclonal antibody. (1)

This indication is approved under accelerated approval based on response rate. Continued approval for this indication may be contingent upon verification and description of clinical benefit in confirmatory trial(s). (1)

Limitations of Use: PEPAXTO is not indicated and is not recommended for use as a conditioning regimen for transplant outside of controlled clinical trials. (1, 5.5)

DOSAGE AND ADMINISTRATION

- Recommended dosage of PEPAXTO is 40 mg intravenously over 30 minutes on Day 1 of each 28-day treatment cycle, in combination with dexamethasone. (2.1)
- See Full Prescribing Information for instructions on preparation and administration. (2.4)

DOSAGE FORMS AND STRENGTHS

For injection: 20 mg melphalan flufenamide as a lyophilized powder in single-dose vial for reconstitution and dilution. (3)

CONTRAINDICATIONS

History of serious hypersensitivity reaction to melphalan flufenamide or melphalan. (4)

WARNINGS AND PRECAUTIONS

- Thrombocytopenia: Monitor platelet counts at baseline, during treatment, and as clinically indicated. Dose delay or dose reduction may be required to allow recovery of platelets. (2.3, 5.1)
- Neutropenia: Monitor neutrophil counts at baseline, during treatment and as clinically indicated. Monitor patients with neutropenia for signs of infection. Dose delay or dose reduction may be required to allow recovery of neutrophils. (2.3, 5.2)
- Anemia: Monitor red blood cell counts at baseline, during treatment, and as clinically indicated. (5.3)
- Infections: Monitor for signs/symptoms of infection and treat promptly. (5.4)
- Increased Risk of Mortality with PEPAXTO at Dosages Higher than the Recommended Dosage: Dosages exceeding the recommended dose for PEPAXTO may be associated with mortality. (1, 5.5, 13.2)
- Secondary Malignancies: Monitor patients long-term for the development of secondary malignancies. (5.6)
- Embryo-Fetal Toxicity: Can cause fetal harm. Advise patients of reproductive potential of the potential risk to a fetus and to use effective contraception. (5.7, 8.1, 8.3)

ADVERSE REACTIONS

Most common adverse reactions (> 20%) are fatigue, nausea, diarrhea, pyrexia and respiratory tract infection. (6.1)

Most common laboratory abnormalities (≥50%) are leukocytes decrease, platelets decrease, lymphocytes decrease, neutrophils decrease, hemoglobin decrease and creatinine increase. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Oncopeptides Inc at 1-866-522-8894 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

USE IN SPECIFIC POPULATIONS

Lactation: Advise not to breastfeed. (8.2)

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

Revised: 2/2021

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1 INDICATIONS AND USAGE

PEPAXTO is indicated in combination with dexamethasone for the treatment of adult patients with relapsed or refractory multiple myeloma who have received at least four prior lines of therapy and whose disease is refractory to at least one proteasome inhibitor, one immunomodulatory agent, and one CD38-directed monoclonal antibody.

This indication is approved under accelerated approval based on response rate. Continued approval for this indication may be contingent upon verification and description of clinical benefit in confirmatory trial(s) [see *Clinical Studies (14)*].

Limitations of Use

PEPAXTO is not indicated and is not recommended for use as a conditioning regimen for transplant outside of controlled clinical trials [see *Warnings and Precautions (5.5)*].

2 DOSAGE AND ADMINISTRATION

2.1 Recommended Dosage

The recommended dosage of PEPAXTO is 40 mg administered intravenously over 30 minutes on Day 1 of each 28-day cycle until disease progression or until unacceptable toxicity. Administer dexamethasone 40 mg orally or intravenously on Days 1, 8, 15 and 22 of each cycle. For patients 75 years of age or older, reduce the dose of dexamethasone to 20 mg. Refer to the prescribing information for dexamethasone for additional dosing information [see *Clinical Studies (14)*].

2.2 Recommended Premedication and Concomitant Medications

Consider providing a serotonin-3 (5-HT₃) receptor antagonist or other antiemetics prior to and during the treatment with PEPAXTO.

2.3 Dosage Modifications for Adverse Reactions

Withhold PEPAXTO if the neutrophil count is less than $1 \times 10^9/L$ or the platelet count is less than $50 \times 10^9/L$.

The recommended dose reductions and dosage modifications for adverse reactions for PEPAXTO are presented in Table 1 and Table 2, respectively.

Table 1: Recommended Dose Reductions for Adverse Reactions of PEPAXTO

Dose Reduction	Dosage*
First	30 mg
Second	20 mg
Subsequent	Permanently discontinue PEPAXTO in patients who are unable to tolerate 20 mg.

* Administered intravenously on Day 1 of each 28-day cycle. For dosage modifications, see Table 2.

Table 2: Recommended Dosage Modifications for Adverse

Reactions of PEPAXTO

Adverse Reaction	Severity	Dosage Modification
Myelosuppression <i>[see Warnings and Precautions (5.1, 5.2)]</i>	Platelet count less than $50 \times 10^9/L$ on an intended PEPAXTO dosing day	<ul style="list-style-type: none"> • Withhold PEPAXTO and monitor platelet count weekly until platelet count is $50 \times 10^9/L$ or greater. • Resume PEPAXTO <ul style="list-style-type: none"> - at same dose if delay is 2 weeks or less. - at 1 dose level lower if delay is more than 2 weeks.
	Absolute neutrophil count less than $1 \times 10^9/L$ on an intended PEPAXTO dosing day	<ul style="list-style-type: none"> • Withhold PEPAXTO and monitor neutrophil count weekly until neutrophil count is $1 \times 10^9/L$ or greater. • Resume PEPAXTO <ul style="list-style-type: none"> - at same dose if delay is 2 weeks or less. - at 1 dose level lower if delay is more than 2 weeks.
	Grade 4 hematological adverse reaction on an intended PEPAXTO dosing day in 2 consecutive cycles	<ul style="list-style-type: none"> • Resume PEPAXTO at 1 dose level lower.
Non-Hematologic Adverse Reaction <i>[see Adverse Reactions (6.1)]</i>	Grade 2	<ul style="list-style-type: none"> • Consider withholding PEPAXTO until resolved to at least Grade 1 or baseline. • Consider resuming PEPAXTO at 1 dose level lower.
	Grade 3 or 4	<ul style="list-style-type: none"> • Withhold PEPAXTO until resolved to at least Grade 1 or baseline. • Resume PEPAXTO at 1 dose level lower as clinically appropriate.

2.4 Preparation and Administration

PEPAXTO is a hazardous drug. Follow applicable special handling and disposal procedures.¹

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Do not use if visibly opaque particles, discoloration or foreign particles are observed.

Reconstitute and dilute PEPAXTO prior to infusion.

Additional agents required for preparation:

- 5% Dextrose Injection, USP (room temperature)
- 250 mL bag of **cold (2°C to 8°C / 36°F to 46°F)** 0.9% Sodium Chloride Injection, USP (refrigerate for at least 4 hours)

Preparation Steps:

Read the complete instructions prior to starting preparation.

Steps 3 to 5 must be completed within 30 minutes.

Reconstitution and dilution steps
Step 1 Determine the dose, the total volume of reconstituted PEPAXTO solution required, and the number of PEPAXTO vials needed. More than one vial may be needed for a full dose. Place PEPAXTO vial(s) at room temperature for at least 30 minutes.
Step 2 Shake the vial(s) vigorously or vortex to disintegrate the lyophilized PEPAXTO powder cake into a loose powder.
Step 3 to 5 must be completed within 30 minutes
Step 3 Aseptically reconstitute each vial with 40 mL of 5% Dextrose Injection, USP to obtain a final concentration of 0.5 mg/mL. Ensure the 5% Dextrose Injection, USP is room temperature (20°C to 25°C / 68°F to 77°F). Shake the vial(s) vigorously until solution is clear. Let the vial(s) stand to allow air bubbles to dissipate to confirm a clear solution.
Step 4 Withdraw 80 mL from a refrigerated (2°C to 8°C / 36°F to 46°F) 250 mL infusion bag of 0.9% Sodium Chloride Injection, USP. Discard the withdrawn 80 mL.
Step 5 Withdraw the required volume of reconstituted solution from the PEPAXTO vial(s) and transfer into an intravenous bag containing 0.9% Sodium Chloride Injection, USP to obtain a final concentration of 0.1 mg/mL to 0.16 mg/mL. Discard any unused portion left in the vial(s). Gently invert the bag to mix the solution. Do not shake. Check that the PEPAXTO solution is clear and colorless to pale yellow. Do not use if solution discoloration or particles are observed.

Storage timelines:

PEPAXTO degrades in solution, especially at room temperature, and the storage timelines for diluted solution should not be exceeded:

For immediate administration:
Infusion of the diluted PEPAXTO solution must begin within 60 minutes of start of reconstitution (step 3).
For delayed administration:
If not used for immediate administration, the diluted PEPAXTO solution should be placed in a refrigerator (2°C to 8°C / 36°F to 46°F) within 30 minutes after initial reconstitution (step 3) and store for up to 6 hours .

Administration:

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Do not use if visibly opaque particles, discoloration or foreign particles are observed.

Administration steps
Step 6 Administer PEPAXTO as a 30-minute intravenous infusion via a central venous access device, for example mediport, PICC or tunneled central venous catheter. If the infusion bag has been stored in a refrigerator, allow to reach to room temperature (20°C to 25°C / 68°F to 77°F). Start infusion within 30 minutes of removing the diluted PEPAXTO solution from the refrigerator.
Step 7 Administer PEPAXTO as an intravenous infusion via a central catheter over 30 minutes.
Step 8 Upon completion of PEPAXTO infusion, flush the central catheter per individual institutional guidelines.

3 DOSAGE FORMS AND STRENGTHS

For Injection: 20 mg melphalan flufenamide as a sterile lyophilized white to off-white powder in a single-dose vial for reconstitution and further dilution.

4 CONTRAINDICATIONS

PEPAXTO is contraindicated in patients with a history of serious hypersensitivity reaction to melphalan flufenamide or melphalan [see *Adverse Reactions (6.1)*].

5 WARNINGS AND PRECAUTIONS

5.1 Thrombocytopenia

Thrombocytopenia was reported in 99% of 157 patients who received PEPAXTO with dexamethasone. Grade 3 thrombocytopenia was reported in 26% and Grade 4 thrombocytopenia was reported in 54% of patients [see *Adverse Reactions (6.1)*]. Thrombocytopenia may lead to hemorrhage. Any Grade hemorrhage was reported in 28% of 157 patients. Grade 3 hemorrhage was reported in 3.2% and Grade 4 hemorrhage was reported in <1% of patients [see *Adverse Reactions (6.1)*].

Grade 3 or 4 thrombocytopenia occurred in 43% of patients during the first cycle, with a median time to onset of 15 days from the first dose.

Monitor platelets at baseline, during treatment, and as clinically indicated. Monitor more frequently during the first two months of treatment with PEPAXTO. Do not administer PEPAXTO if the platelet count is less than $50 \times 10^9/L$. Withhold PEPAXTO until platelet count $50 \times 10^9/L$ or greater and resume at same or reduced dose based on duration of interruption. Adjust dose and/or dose schedule based on signs and symptoms of bleeding [see *Dosage and Administration (2.3)*].

5.2 Neutropenia

Neutropenia was reported in 95% of 157 patients who received PEPAXTO with dexamethasone. Grade 3 neutropenia was reported in 41% and Grade 4 neutropenia was reported in 40% of patients. Febrile neutropenia was reported in 6% of patients [see *Adverse Reactions (6.1)*]. Neutropenia may lead to infection.

Grade 3 or 4 neutropenia occurred in 50% during the first cycle, with a median time to onset of 15 days from the first dose.

Monitor neutrophil counts at baseline, during treatment, and as clinically indicated. Monitor more frequently during the first two months of treatment with PEPAXTO. Do not administer PEPAXTO if absolute neutrophil count less than $1 \times 10^9/L$. Withhold PEPAXTO until absolute neutrophil count is $1 \times 10^9/L$ or greater and resume at same or reduced dose based on duration of interruption [see *Dosage and Administration (2.3)*]. Consider leukocyte growth factor as clinically appropriate.

5.3 Anemia

Anemia was reported in 84% of 157 patients who received PEPAXTO with dexamethasone. Grade 3 anemia was reported in 50% of 157 patients [see *Adverse Reactions (6.1)*].

Monitor red blood cell counts at baseline, during treatment, and as clinically indicated. Monitor more frequently during the first two months of treatment with PEPAXTO. Treat anemia as clinically indicated and as per standard guidelines. Dosage modification and dose delay of PEPAXTO may be required to allow for recovery of red blood cells.

5.4 Infections

Fatal infections were reported in <1% of 157 patients who received PEPAXTO with dexamethasone. Any Grade infection was reported in 58% of 157 patients who received PEPAXTO and dexamethasone. Grade 3 infections were reported in 20% and Grade 4 infection was reported in 1.9% of patients. Respiratory tract infection occurred in 24% (Grade ≥ 3 in 5%), pneumonia in 13% (Grade ≥ 3 in 11%), and sepsis in 3.8% (Grade ≥ 3

in 3.2%) of patients [see *Adverse Reactions (6.1)*]. Consider antimicrobials as clinically appropriate.

5.5 Increased Risk of Mortality with PEPAXTO at Dosages Higher than the Recommended Dosage

A nonclinical safety study in dogs with melphalan flufenamide at dosages exceeding the recommended dose for relapsed or refractory multiple myeloma was associated with mortality [see *Nonclinical Toxicology (13.2)*]. There is limited clinical experience of PEPAXTO at dosages higher than recommended. The safety and efficacy of PEPAXTO has not been established for use as a conditioning regimen in patients receiving transplant.

5.6 Secondary Malignancies

Secondary malignancies such as myelodysplastic syndromes or acute leukemia have occurred in patients with multiple myeloma who have received PEPAXTO. Monitor patients long-term for the development of secondary malignancies.

5.7 Embryo-Fetal Toxicity

Based on its mechanism of action, PEPAXTO can cause fetal harm when administered to a pregnant woman because it is genotoxic and targets actively dividing cells. Advise pregnant women of the potential risk to a fetus. Advise females of reproductive potential to use effective contraception during treatment with PEPAXTO and for 6 months after the last dose. Advise males with female partners of reproductive potential to use effective contraception during treatment with PEPAXTO and for 3 months after the last dose [see *Use In Specific Populations (8.1, 8.3)*].

6 ADVERSE REACTIONS

The following clinically significant adverse reactions are described elsewhere in the labeling:

- Thrombocytopenia [see *Warnings and Precautions (5.1)*].
- Neutropenia [see *Warnings and Precautions (5.2)*].
- Anemia [see *Warnings and Precautions (5.3)*].
- Infections [see *Warnings and Precautions (5.4)*].

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.

Relapsed Refractory Multiple Myeloma (RRMM)

The safety of PEPAXTO was evaluated in HORIZON [see *Clinical Studies (14)*]. Patients received PEPAXTO 40 mg intravenously on Day 1 of each 28-day cycle, in combination with dexamethasone 40 mg orally (or 20 mg for patients 75 years and older) on Days 1, 8, 15 and 22 of each cycle (N=157). Patients were enrolled if they had absolute neutrophil count of $1 \times 10^9/L$ or higher and platelet count of $75 \times 10^9/L$ or greater. Among patients who received PEPAXTO, 29% were exposed for 6 months or longer and

6% were exposed for greater than one year.

Serious adverse reactions occurred in 49% of patients who received PEPAXTO. Serious adverse reactions in >3% of patients included pneumonia (10%), respiratory tract infection (6%), thrombocytopenia (5%), febrile neutropenia (5%) and sepsis (3.2%). Fatal adverse reactions occurred in 10 patients (6%) who received PEPAXTO, where general physical health deterioration (1.9%) and respiratory failure (1.3%) represented more than 1%.

Permanent discontinuation of PEPAXTO due to an adverse reaction occurred in 22% of patients. Adverse reactions which resulted in permanent discontinuation of PEPAXTO in >3% of patients included thrombocytopenia (11%).

Dosage interruptions of PEPAXTO due to an adverse reaction occurred in 62% of patients. The adverse reactions which resulted in dosage interruption of PEPAXTO in >3% of patients included thrombocytopenia (43%), neutropenia (29%), anemia (10%), respiratory tract infection (7%), leukopenia (6%) and pyrexia (4.5%).

Dose reductions of PEPAXTO due to an adverse reaction occurred in 27% of patients. Adverse reactions which resulted in dose reductions of PEPAXTO in >3% patients included thrombocytopenia (22%) and neutropenia (6%).

The most common adverse reactions ($\geq 20\%$) were fatigue, nausea, diarrhea, pyrexia and respiratory tract infection. The most common laboratory abnormalities ($\geq 50\%$) were leukocytes decreased, platelets decreased, lymphocytes decreased, neutrophils decreased, hemoglobin decreased and creatinine increased.

Table 3 summarizes the adverse reactions in HORIZON.

Table 3: Adverse reactions ($\geq 10\%$) in Patients with RRMM Who Received PEPAXTO with Dexamethasone in HORIZON

Adverse Reaction	PEPAXTO with Dexamethasone (N=157)	
	All Grades (%)	Grade 3 or 4 (%)
<i>General disorders and administration site disorders</i>		
Fatigue*	55	6
Pyrexia†	24	1.9
Edema peripheral†	14	1.3
<i>Gastrointestinal disorders</i>		
Nausea†	32	0.6
Diarrhea	27	0
Constipation†	15	0.6
Vomiting	13	0
<i>Infections</i>		
Respiratory tract infection†,‡	24	5
Pneumonia§	13	11
<i>Respiratory, thoracic and</i>		

<i>mediastinal disorders</i>		
Cough [†]	17	0
Dyspnea [†]	11	1.3
Dyspnea exertional	10	0
<i>Metabolism and nutrition disorders</i>		
Decreased appetite [†]	14	0.6
Hypokalemia [†]	14	1.3
Hypocalcemia [†]	10	0.6
<i>Nervous system disorders</i>		
Headache	13	0
Dizziness	11	0
<i>Musculoskeletal and connective tissue disorders</i>		
Bone pain [†]	13	1.9
Pain in extremity [†]	13	1.9
Back pain [†]	12	0.6
Arthralgia	10	0
<i>Psychiatric disorders</i>		
Insomnia [†]	11	0.6

* Fatigue includes fatigue and asthenia

† No Grade 4 adverse reactions occurred

‡ Respiratory tract infection includes upper respiratory tract infection, lower respiratory tract infection, respiratory tract infection and respiratory tract infection viral

§ Pneumonia includes pneumonia, pneumocystis jirovecii pneumonia and pneumonia viral

Clinically relevant adverse reactions in <10% of patients who received PEPAXTO in combination with dexamethasone (N=157) included:

Allergic conditions: hypersensitivity reaction (7%)

Blood and lymphatic system disorders: febrile neutropenia (6%)

Infections: sepsis (3.8%)

Hemorrhages: Grade 3 or 4 hemorrhages (3.8%)

Table 4 summarizes the laboratory abnormalities in HORIZON.

Table 4: Laboratory Abnormalities (≥50%) That Worsened from Baseline in Patients in HORIZON

Laboratory Abnormality	PEPAXTO with Dexamethasone*	
	All Grades[†] (%)	Grade 3- 4[‡] (%)
Leukocytes decrease	99	88
Platelets decrease	99	80
Lymphocytes decrease	97	95
Neutrophils decrease	95	82
Hemoglobin decrease	84	50

Creatinine increase	68	1 [§]
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* Denominators for percentages are the number of patients with assessments at baseline and post-baseline (N=157 for all abnormalities)

† Patients with any worsening grade

‡ Patients with worsening to Grade 3 or 4, respectively

§ No Grade 4 laboratory abnormality occurred

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

Based on its mechanism of action [see *Clinical Pharmacology (12.1)*], PEPAXTO can cause fetal harm when administered to a pregnant woman. There are no available data on PEPAXTO use in pregnant women to evaluate for a drug-associated risk. PEPAXTO is a genotoxic drug [see *Nonclinical Toxicology (13.1)*]. Advise pregnant women 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

Animal reproductive or developmental toxicity studies were not conducted with PEPAXTO. Melphalan flufenamide is genotoxic and was toxic to actively dividing cells in animal studies and thus it has the potential to cause teratogenicity and embryo-fetal lethality.

8.2 Lactation

Risk Summary

There is no data on the presence of melphalan flufenamide or its metabolites in human breast milk, or the effects on the breastfed child or on milk production. Because of the potential for serious adverse reactions in the breastfed child, advise women not to breastfeed during treatment with PEPAXTO and for 1 week after the last dose.

8.3 Females and Males of Reproductive Potential

PEPAXTO can cause fetal harm when administered to a pregnant woman [see *Use in Specific Populations (8.1)*].

Pregnancy Testing

Verify pregnancy status in females of reproductive potential prior to initiating PEPAXTO.

Contraception

Females

Advise females of reproductive potential to use effective contraception during treatment with PEPAXTO and for 6 months after the last dose.

Males

Based on genotoxicity findings, advise males with female partners of reproductive potential to use effective contraception during treatment with PEPAXTO and for 3 months after the last dose [see *Nonclinical Toxicology (13.1)*].

Infertility

Females

PEPAXTO can cause amenorrhea in premenopausal women and result in infertility.

Males

Based on findings of melphalan flufenamide in animals, PEPAXTO may impair male fertility [see *Nonclinical Toxicology (13.1)*]. Alkylating drugs, such as PEPAXTO, can also cause irreversible testicular suppression in patients.

8.4 Pediatric Use

The safety and effectiveness of PEPAXTO have not been established in pediatric patients.

8.5 Geriatric Use

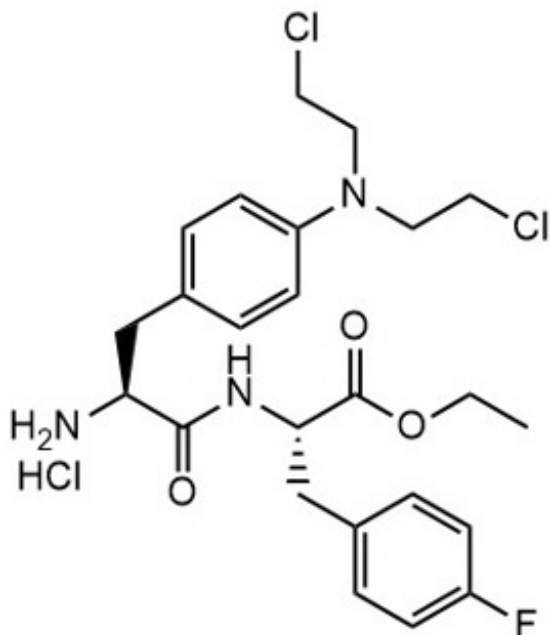
Of the 157 patients with RRMM who received PEPAXTO, 50% were 65 years and older, while 16% were 75 years and older. No overall differences in safety were observed between these patients and younger patients. Clinical studies of PEPAXTO in patients with RRMM did not include sufficient numbers of patients 65 years of age and older to determine if they respond differently from younger adult patients.

8.6 Renal Impairment

No dose adjustment of PEPAXTO is recommended in patients with creatinine clearance (CLcr) 45 to 89 mL/min calculated using Cockcroft-Gault equation [see *Clinical Pharmacology (12.3)*]. PEPAXTO has not been studied in patients with CLcr 15 to 44 mL/min.

11 DESCRIPTION

Melphalan flufenamide is an alkylating drug. The chemical name is Ethyl (2S)-2-[[[(2S)-2-amino-3-[4-[bis(2-chloroethyl)amino]phenyl]propanoyl]amino]-3-(4-fluorophenyl)propanoate hydrochloride and the molecular weight is 498.4 as free base and 534.9 as the hydrochloride salt. The structural formula is:



Melphalan flufenamide hydrochloride is soluble in most organic solvents, while sparsely soluble in aqueous solutions. The pKa value is 7.13.

PEPAXTO for injection is supplied as a sterile, white to off-white lyophilized powder in a single-dose vial for intravenous use. Each vial contains 20 mg melphalan flufenamide (equivalent to 21.48 mg melphalan flufenamide hydrochloride) and 1,000 mg sucrose.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Melphalan flufenamide is a peptide conjugated alkylating drug. Due to its lipophilicity, melphalan flufenamide is passively distributed into cells and thereafter enzymatically hydrolyzed to melphalan. Similar to other nitrogen mustard drugs, cross-linking of DNA is involved in the antitumor activity of melphalan flufenamide. In cellular assays, melphalan flufenamide inhibited proliferation and induced apoptosis of hematopoietic and solid tumor cells. Additionally, melphalan flufenamide showed synergistic cytotoxicity with dexamethasone in melphalan resistant and non-resistant multiple myeloma cell lines.

12.2 Pharmacodynamics

The exposure-response relationship and time course of pharmacodynamic response for the safety and effectiveness of PEPAXTO have not been fully characterized.

Cardiac Electrophysiology

The effect of PEPAXTO on QT interval has not been fully characterized.

12.3 Pharmacokinetics

Melphalan flufenamide peak plasma concentrations were reached during the 30-minute infusion. Peak plasma concentrations of the active metabolite melphalan were reached 4 to 15 minutes after the end of infusion of PEPAXTO 40 mg. Following PEPAXTO 40 mg, the mean (CV%) C_{max} was 432 ng/mL (30%) and AUC_{0-1NF} was 3,143 $\mu\text{g/mL}\cdot\text{hr}$ (28%)

for melphalan after a single dose. The mean (CV%) C_{max} was 419 ng/mL (33%) and AUC_{0-INF} was 2,933 $\mu\text{g/mL}\cdot\text{hr}$ (29%) for melphalan at steady-state.

Distribution

In vivo the disappearance of melphalan flufenamide from plasma is rapid and is attributed to distribution to peripheral tissues with no late redistribution back to plasma.

The mean (CV%) volume of distribution was 35 L (71%) for melphalan flufenamide and 76 L (32%) for melphalan after a single dose.

Elimination

After the end of infusion of PEPAXTO 40 mg, the mean (CV%) elimination half-life of melphalan flufenamide is 2.1 minutes (34%). The mean (CV%) elimination half-life of melphalan is 70 minutes (21%). The mean (CV%) clearance of melphalan flufenamide and melphalan is 692 L/hr (49%) and 23 L/hr (23%), respectively, at the recommended dosage of PEPAXTO 40 mg.

Metabolism

Melphalan flufenamide is metabolized in tissues to desethyl-melphalan flufenamide and melphalan. Melphalan is metabolized primarily by spontaneous hydrolysis to monohydroxy-melphalan and dihydroxy-melphalan.

Specific Populations

Higher melphalan exposures were observed in patients with lower body surface area. No clinically meaningful differences in the PK of melphalan were observed based on age (35 to 85 years old), renal impairment (CL_{cr} 45 to 89 mL/min) and mild hepatic impairment (total bilirubin \leq ULN and $AST >$ ULN, or total bilirubin 1 to 1.5 \times ULN and any AST).

The effect of sex, race/ethnicity, moderate to severe hepatic impairment (total bilirubin $>1.5 \times$ ULN and any AST), and renal impairment (CL_{cr} 15 to 44 mL/min) on melphalan flufenamide and melphalan PK is unknown.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

No carcinogenicity studies have been conducted with melphalan flufenamide.

PEPAXTO is genotoxic. In studies conducted in vitro, melphalan flufenamide caused irreversible DNA damage.

Repeat-dose toxicity studies with melphalan flufenamide in animals showed adverse effects on male reproductive organs. Melphalan flufenamide was administered intravenously to rats at 20, 40, or 55 mg/m^2 , and to dogs at 0.45 or 0.90 mg/kg (9 or 18 mg/m^2) every 21 days for two or three doses. Decreased testes weights and depletion of germ cells were observed in both species, and epididymal oligospermia was observed in dogs. Adverse effects on male reproductive organs were observed in dogs at dose levels less than the recommended clinical dose of 40 mg. The reversibility of adverse effects on male reproductive organs was not assessed.

13.2 Animal Toxicology and/or Pharmacology

Dogs were intravenously administered a single dose of melphalan flufenamide (17.5 mg/kg) or an equimolar dose of melphalan; these dose levels were representative of dosages needed for myeloablation. Increased mortality was observed in dogs administered melphalan flufenamide despite similar melphalan exposure in animals administered melphalan flufenamide or melphalan.

14 CLINICAL STUDIES

The efficacy of PEPAXTO in combination with dexamethasone was evaluated in HORIZON [NCT02963493], a multicenter, single-arm trial. Eligible patients were required to have relapsed or refractory multiple myeloma. Patients received PEPAXTO 40 mg intravenously on Day 1 and dexamethasone 40 mg orally (20 mg for patients ≥ 75 years of age) on Day 1, 8, 15 and 22 of each 28-day cycle until disease progression or unacceptable toxicity.

A total of 157 patients accepting a central venous catheter and with estimated creatinine clearance by Cockcroft-Gault formula ≥ 45 mL/min were enrolled. Patients with primary refractory disease (i.e. never responded with at least minimal response to any prior treatment) were excluded. Ninety seven patients had received four or more prior lines of therapies and were refractory to at least one proteasome inhibitor, at least one immunomodulatory agent and a CD38-directed monoclonal antibody. The median age was 65 years (range: 35 to 86 years); 58% were male, 87% were White and 6% were Black or African American. Disease characteristics in these 97 patients are summarized in Table 5.

The major efficacy outcome measure was overall response rate (ORR) and Duration of Response (DoR) assessed according to the International Myeloma Working Group (IMWG) Criteria by investigators. Efficacy results in the 97 patients are provided in Table 6. The median time to first response was 2.1 months (range: 1.0 to 6.1 months).

Table 5: Disease Characteristics (HORIZON)

Parameter	PEPAXTO with Dexamethasone (N=97)
Years from diagnosis to start of PEPAXTO, median (range)	6.4 (2.1 to 24.6)
Prior treatment regimens, median (range)	6 (4 to 12)
Documented refractory status, (%)	
Lenalidomide	94
Pomalidomide	92
Bortezomib	74
Carfilzomib	63
Daratumumab	93
Alkylator refractory, (%)	75
Previous stem cell transplant, (%)	70
International Staging System at Baseline, (%)	
I	30

II	32
III	34
Missing/Unknown	4
High-risk cytogenetics*, (%)	33
Extramedullary disease (EMD), (%)	41

* del(17p), t(4;14),t(14;16), gain (1q) and t(14;20) at study entry

Table 6: Efficacy Results (HORIZON)

	PEPAXTO with Dexamethasone (N=97)
Overall response rate (ORR), n (%) (95% CI)	23 (23.7) (15.7, 33.4)
Stringent complete response (sCR)	0
Complete Response (CR)	0
Very good partial response (VGPR), n (%)	9 (9.3)
Partial response (PR), n (%)	14 (14.4)
Median duration of response in months (95% CI)	4.2 (3.2, 7.6)

15 REFERENCES

1. "OSHA Hazardous Drugs." OSHA.
<http://www.osha.gov/SLTC/hazardousdrugs/index.html>

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied

PEPAXTO is a white to off-white lyophilized powder for reconstitution (after reconstitution the solution is clear and colorless to light yellow) supplied in a 50 mL single dose vial containing 20 mg melphalan flufenamide. Each 20 mg vial is packaged in a single carton (NDC 73657-020-01).

The vial stopper is not manufactured with natural rubber latex.

Storage

Store refrigerated at 2°C to 8°C (36°F to 46°F) and protect from light. Retain in original carton until use.

Handling and Disposal

PEPAXTO is a hazardous drug. Follow special handling and disposal procedures.¹ All materials that have been utilized for dilution and administration, including any reconstituted solution made over 30 minutes prior, should be disposed of according to standard procedures for hazardous drugs.

17 PATIENT COUNSELING INFORMATION

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

Thrombocytopenia, Neutropenia and Anemia

- Advise patients that PEPAXTO can cause myelosuppression. Advise patients to immediately report signs or symptoms of thrombocytopenia (bleeding and easy bruising), neutropenia (symptoms of infection, such as fever, chills, cough, pain, or burning during urination) and anemia (fatigue and shortness of breath) to their healthcare provider.
- Advise patients that complete blood counts will be monitored at baseline, during treatment, and as clinically indicated [*see Warnings and Precautions (5.1, 5.2, 5.3)*].

Infections

Advise patients that PEPAXTO can cause infections. Instruct patients to immediately report new or worsening signs or symptoms (e.g., chills, fever) of infection to their healthcare provider [*see Warnings and Precautions (5.4)*].

Secondary Malignancies

Advise patients on the risk of second primary malignancies [*see Warnings and Precautions (5.6)*].

Embryo-Fetal Toxicity

- Advise pregnant women and females of reproductive potential of the potential risk to a fetus. Advise females of reproductive potential to inform their healthcare provider of a known or suspected pregnancy [*see Warnings and Precautions (5.7) and Use in Specific Populations (8.1)*].
- Advise females of reproductive potential to use effective contraception during treatment with PEPAXTO and 6 months after the last dose [*see Use in Specific Populations (8.3)*].
- Advise males with female partners of reproductive potential to use effective contraception during treatment with PEPAXTO and for 3 months after the last dose [*see Use in Specific Populations (8.3)*].

Lactation

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

Manufactured for: Oncopeptides AB (publ), Stockholm, Sweden.

Distributed by: Oncopeptides Inc. 200 Fifth Avenue, Suite #1030 Waltham, MA 02451, USA

PEPAXTO is a registered trademark of Oncopeptides AB (publ)

PATIENT INFORMATION

PEPAXTO (peh-PAX-toe)

(melphalan flufenamide)

for injection, for intravenous use

What is PEPAXTO?

PEPAXTO is a prescription medicine used in combination with the medicine

dexamethasone to treat adults with multiple myeloma who did not respond to or stopped responding to at least four prior medicines including at least one proteasome inhibitor, one immunomodulatory agent and one CD38-directed antibody.

PEPAXTO is not for use to prepare for transplant.

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

Do not receive PEPAXTO if you have a history of a severe allergic reaction to melphalan flufenamide or melphalan.

Before receiving PEPAXTO, tell your healthcare provider about all of your medical conditions, including if you:

- have an infection
- are pregnant or plan to become pregnant. PEPAXTO may harm your unborn baby.

Females who are able to become pregnant:

- Your healthcare provider will check to see if you are pregnant before you start treatment with PEPAXTO.
- You should use an effective method of birth control (contraception) during treatment and for 6 months after the last dose of PEPAXTO.
- Tell your healthcare provider right away if you become pregnant or think you may be pregnant during treatment with PEPAXTO.

Males with female partners who are able to become pregnant:

- You should use an effective method of birth control (contraception) during treatment and for 3 months after the last dose of PEPAXTO.
- Talk to your healthcare provider about effective methods of birth control that you can use during this time.
- are breastfeeding or plan to breastfeed. It is not known if PEPAXTO passes into breast milk. Do not breastfeed during treatment with and for 1 week after the last dose of PEPAXTO.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

How will I receive PEPAXTO?

- PEPAXTO is given to you by your healthcare provider into your vein through intravenous (IV) infusion over 30 minutes.
- PEPAXTO is usually given 1 time every 28 days.
- Your healthcare provider will decide how many treatments you need.
- Your healthcare provider will do blood tests before and during your treatment with PEPAXTO to check for side effects.
- Your healthcare provider may prescribe medicines to help prevent nausea before your infusion and during treatment with PEPAXTO.

What are the possible side effects of PEPAXTO?

PEPAXTO may cause serious side effects, including:

- **Low blood cell counts** are common with PEPAXTO and can be serious. Your healthcare provider will do blood tests as needed to check your blood cell counts during your treatment with PEPAXTO.
 - **Low platelet counts:** Tell your healthcare provider right away if you have a bleeding or bruising under the skin.
 - **Low red blood cell counts:** Tell your healthcare provider if you are feeling weak, tired or you get tired easily, you look pale, or if you feel short of breath.
 - **Low white blood cell counts:** A low white blood cell count increases the risk of

infections.

- **Infections.** PEPAXTO can cause infections that have led to death. **Tell your healthcare provider right away if you develop new or worsening signs or symptoms of infection such as fever, chills, cough, pain, or burning during urination during treatment with PEPAXTO.**
- **Secondary cancers.** New cancers such as myelodysplastic syndromes or acute leukemia have happened in people with multiple myeloma who have received PEPAXTO. Your healthcare provider will monitor you for new cancers.

Your healthcare provider may change your dose of PEPAXTO, stop your treatment for a period of time, or completely stop your treatment if you have certain side effects. PEPAXTO may cause fertility problems in males and females, which may affect your ability to have children. Talk with your healthcare provider if you have concerns about fertility.

The most common side effects of PEPAXTO include, low blood cell counts, fatigue, nausea, diarrhea, fever, and cold-like symptoms (respiratory tract infection).

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

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

General information about the safe and effective use of PEPAXTO.

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. You can ask your healthcare provider or pharmacist for information about PEPAXTO that is written for health professionals.

What are the ingredients in PEPAXTO?

Active ingredient: melphalan flufenamide

Inactive ingredient: sucrose

Manufactured for Oncopeptides AB (publ), Stockholm, Sweden.

Marketed and distributed by Oncopeptides Inc., 200 Fifth Avenue, Suite #1030 Waltham, MA 02451, USA.

PEPAXTO is a registered trademark of Oncopeptides AB (publ).

For more information, go to www.PEPAXTO.com or call 1-866-522-8894.

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

Issued:
02/2021

PRINCIPAL DISPLAY PANEL - 20 mg Vial Label

NDC 73657-020-01

Rx only

Pepaxto®

(melphalan flufenamide)

for injection

20 mg/vial

For Intravenous Infusion Only.

Reconstitute and Dilute prior to administration.

Discard unused portion.

CAUTION: Hazardous Agent

One single-dose vial


618133

Each vial contains 20 mg melphalan flufenamide and 1,000 mg sucrose.

Store refrigerated at 2°C - 8°C (36°F - 46°F) in the carton to protect from light. Do not freeze.

Dosage: See Prescribing Information.

Manufactured for Oncopeptides AB (publ)

 oncopeptides

NDC 73657-020-01 Rx only

Pepaxto[®]
(melphalan flufenamide)
for injection

20 mg/vial



For Intravenous Infusion Only.
Reconstitute and Dilute prior to administration.

Discard unused portion.

CAUTION: Hazardous Agent
One single-dose vial

618133

EXP YYYY-MMM
Lot



PEPAXTO

melphalan flufenamide injection, powder, lyophilized, for solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:73657-020
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
melphalan flufenamide hydrochloride (UNII: 3412470A0V) (melphalan flufenamide - UNII:F70C5K4786)	melphalan flufenamide	20 mg in 50 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73657-020-01	1 in 1 CARTON	02/26/2021	
1		50 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA214383	02/26/2021	

Labeler - Oncopeptides, AB (632402728)

Establishment

Name	Address	ID/FEI	Business Operations
Magle Chemoswed AB		559533158	API MANUFACTURE(73657-020)

Establishment

Name	Address	ID/FEI	Business Operations
Cenexi-Laboratoires Thissen SA (Cenexi LT)		370088959	MANUFACTURE(73657-020) , ANALYSIS(73657-020)

Establishment

Name	Address	ID/FEI	Business Operations
Apotek Produktion & Laboratorier AB (APL)		632474078	ANALYSIS(73657-020)

Establishment

Name	Address	ID/FEI	Business Operations
Eurofins Biopharma Product Testing, Denmark A/S (Eurofins)		311900950	ANALYSIS(73657-020)

Establishment

Name	Address	ID/FEI	Business Operations
Solvias AG (Solvias)		480739627	ANALYSIS(73657-020)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmascience Inc.		202657094	ANALYSIS(73657-020)

Establishment

Name	Address	ID/FEI	Business Operations
AndersonBrecon Inc.; "PCI of Illinois"		098908572	LABEL(73657-020) , PACK(73657-020)

Establishment

Name	Address	ID/FEI	Business Operations
West Pharmaceutical Services, Inc.		002330983	PACK(73657-020)