



Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2021 Introduction Type: New Item Final Version Date: 11/16/2023

PRODUCT INFORMATION **SPECIAL HANDLING AND STORAGE REQUIREMENTS***

Company Name: Camber Pharmaceuticals, Inc. **Application:** ANDA
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 217330
Medical Device Class, if applicable: _____
DUNS: 11-856-3719
Proprietary Name (if Applicable) and Established Name: Famotidine Powder for Oral Suspension, USP 40 mg/5 mL
Selling Unit NDC: 31722-063-31 **Unit of Use NDC:** 31722-063-31 **UPC:** 331722063319
UDI _____ **CVX Code:** _____ **MXV Code:** _____
Description: Famotidine Powder for Oral Suspension, USP 40 mg/5 mL
 NOTE - When reconstituted as directed, famotidine for oral suspension is a white to off-white homogeneous suspension.
Active Ingredient(s): Famotidine
URL for Additional Product Information: www.camberpharma.com
Address: 1031 Centennial Ave (and) 800 Centennial Ave, Suite 1 **Address 2:** _____
City: Piscataway **State:** NJ **Zip:** 08854
Key Contact: Customer Service **Email:** customerservice@camberpharma.com
Phone Number: 1-866-827-3647 **Fax:** 732-562-8788
Product Therapeutic Classification: Histamine H2-receptor antagonist

a. Temperature – Indicate the USP temperature range for this product.
 Temperature Range: Controlled Room – between 20 and 25 C (68° – 77° F)
 Other Temperature Range Requirement (write in): _____
 Notes: Excursions permitted to 15° to 30°C (59° to 86°F)
 Protect from freezing. Discard unused reconstituted suspension after 30 days
 Is this product to be shipped to customers on ice? No
 Is this product to be shipped to customers on dry ice? No
b. Contact for temperature excursion questions:
Name: Soma Raju
Number: 732-529-0423
Group E-mail: somaraju@heterousa.com
c. Special regulations for product in any states? No
 Special returns requirements for this product? No
d. Store product (unit of sale) upright? No
 Protect product (unit of sale) from light? No
e. Shelf life: 24 Months
 Initial shelf life at launch (if different): _____ Months

ADDITIONAL PRODUCT INFORMATION **PRODUCT DESCRIPTION INFORMATION**

The product is a legend device? <input type="checkbox"/> No	Is the Product... Direct-Ship Only <input type="checkbox"/>	Size: 50mL (before reconstitution)
if yes, enter class # _____	Is the Product... Unit of Use <input type="checkbox"/>	
a product kit? <input type="checkbox"/> No	Orphan Drug Status <input type="checkbox"/>	Strength: 40 mg/5 mL
if yes, list NDCs of component parts reverse numbered? _____	FDA Approval Status _____	Dosage Form: Powder for oral suspension
co-licensed? <input type="checkbox"/> No	Allergens Present _____	Product Shape: N/A
latex-free? <input type="checkbox"/> Yes	Corn, Alcohol, Sugar <input type="checkbox"/>	Product Color: White to off-white (see note)
preservative-free? <input type="checkbox"/> No	Country of Origin: India <input type="checkbox"/>	Product Imprint: N/A
correctional institution block? <input type="checkbox"/> No	Is this product covered under the Trade Agreements Act (TAA)? <input type="checkbox"/> No	
opioid? <input type="checkbox"/> No		
Cannabinoid? <input type="checkbox"/> No		
If Unit Dose, is item bar coded to unit dose for hospital scanning? <input type="checkbox"/>		
If Unit Dose, indicate NDC here: _____		

ORDER INFORMATION

Unit of Sale	What is the NDC selling unit?
<input checked="" type="checkbox"/> Bottle	1 Bottle of 50 mL Powder for Oral Suspension
<input type="checkbox"/> Box/Carton	(Write-in, e.g. 1 Box of 10 Vials)
<input type="checkbox"/> Ampule	
<input type="checkbox"/> Glass	Minimum order quantity? <input type="checkbox"/> Yes
<input type="checkbox"/> Tube	
<input type="checkbox"/> Vial Liquid Sgl	
<input type="checkbox"/> Vial Liquid Multi	If Yes, how many of which package type?
<input type="checkbox"/> Vial Powder Sgl	<input type="checkbox"/> 24 Each
<input type="checkbox"/> Vial Power Multi	<input type="checkbox"/> Inner/Carton/Pack
<input type="checkbox"/> Other: Write In _____	<input type="checkbox"/> Case

FOR GENERIC DRUG PRODUCTS

I. Orange Book Rating: AB Authorized Generic *If Authorized Generic, other section fields are not applicable
II. Generic Equivalent to What Brand?: Pepcid

PHARMACY ORDER / BILL UNIT

Rec. sell unit to customer? _____ **Rx billing unit to pharmacy:** _____
 (Write-in, e.g. 1 Vial) Each
 Gram
 Milliliter

DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION

Does supplier meet DSCSA definition of manufacturer? Yes No **GLN:** 0331722000000
Is product exempt from DSCSA? No **GCP:** _____
If yes, select exemption: _____
Other exemption - Write in: _____
Is product repackaged? No **If yes, was original product purchased direct from mfr?** Yes
Is product sold by manufacturer's exclusive distributor? Yes **Provide source manufacturer for repackaged product** _____
Has FDA granted waiver/exception/exemption for product? No
If yes, attach documentation from FDA. _____

ITEM AND PACKING INFORMATION

Item/Each:	Weight Lbs.	Dimensions (US msmts.)			Volume (Cube)	Saleable # Pieces
		Depth	Width	Height		
Item/Each:	0.125	2	2	4	16	1
Box/Carton/Bundle/Inner Pack:						
Case:	3.4	12.5	8.5	5.25	557.81	24
Pallet:						

GTIN AND HIBCC PRODUCT INFORMATION

Saleable Unit of Measure	Saleable Quantity	HIBCC	GTIN-14	Unit of Use GTIN-14
<input checked="" type="checkbox"/> Item/Each	1		00331722063319	00331722063319
<input type="checkbox"/> Box/Carton/Bundle/Inner Pack				
<input checked="" type="checkbox"/> Case	24		20331722063313	
<input type="checkbox"/> Pallet				

COST INFORMATION **WHOLESALE USE ONLY:**

Regular Cost _____ **Vendor #:** _____
Invoice Cost (WAC) (\$) \$22.00 **Whsl. Code #:** _____
As of date: 11/15/2023 **Fineline Code:** _____



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MATERIAL HAZARD CLASSIFICATION and TRANSPORTATION

Is this product (check all that apply):

- a. Cytotoxic? No
- b. CA Prop. 65 Carcinogen or Reproductive Toxicant?
 - Is the product a CA Prop 65 carcinogen? No
 - Is the product a CA Prop 65 reproductive toxicant? No
 - Does the product label bear a CA Prop 65 warning? No

- c. Contact Hazard? No
- d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) No
- e. Does the product contain DEHP? No

Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS)

- a. UN/Identification Number
- b. Proper Shipping Name
- c. DOT Hazard Class
- d. Packing Group
- e. Inhalation Hazard? No

Is this product regulated for shipment by IATA? (if yes, answer a-e below and provide SDS)

- a. UN/Identification Number
- b. Proper Shipping Name
- c. DOT Hazard Class
- d. Packing Group
- e. Inhalation Hazard? No

Is the product restricted for air shipment? If so, indicate restriction:

- Passenger No
- Cargo
- Passenger & Cargo

Is this a reportable quantity? No

RQ Threshold:

Is this a marine pollutant? No

Is this product shipped utilizing an authorized DOT exception or Special Permit?

- No (if yes, identify method below)
- Limited Quantity
- Consumer Commodity, ORM-D
- Small Quantity (49 CFR 173.4)
- Special Permit; DOT-SP
- Special Provision (listed in Column 7 of 49 CFR 172.101); SP#

ADD'L STORAGE INFORMATION

Is the Product...

- Controlled Substance? No Yes Controlled Substance Code
- Controlled by State(s)? No Yes Listed Chemical (List I or II) No Yes
- ARCOS Reportable? No Yes If yes, indicate which:
- Schedule No. Is it a scheduled listed chemical product?: No Yes

CLASS OF TRADE RESTRICTION:

- No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes No
- Restricted to retail pharmacy only: No Yes
- Restricted to hospital, clinics, and physician offices only: No Yes
- Restricted from US territories? (explain in comments) No Yes

Comments:

SDS Hazard Classification

- Organic Corrosive
- Inorganic Oxidizer
- Steroid/Androgen Contact Hazard

Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No

Is the product a NIOSH hazardous drug? If yes, indicate which: No

Hazardous Waste Identification

EPA Hazardous Waste Code: Waste Characteristics:

REMS or REGISTRY RESTRICTIONS

Is there a REMS on this product? No Yes Website URL:

Med Guide Required No Yes Limited Distribution Requirement No Yes Comments / Details: (For example, iPledge program?)

REMS: No Yes REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: No Yes Wholesale distributor support: No Yes Provider Name: DEA #: Site Enrollment Number assigned by Supplier: NCPDP#: NPI #:

Comments

Registry: No Yes Registry Program Contact Name: Phone: Comments

RETURN INSTRUCTIONS

Contact tel. # if product received damaged: 1-866-827-3647

Is product returnable for credit: Yes No

URL/Link to returns policy: contact - customerservice@camberpharma.com

Special regulations or returns requirements for this product in certain states? No Yes

If so, which states? Other requirements? Comments?

MISCELLANEOUS NOTES and/or Image of Product Barcode:

