

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

Version 2021						Introduction Type:	New Item		x Final Version			Date:	6/1/	/2022
			PRODUCT INFORMA	TION					SPECIAL HAN	IDLING AND STOR	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA					ANDA	a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 215767							Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)							
Medical Device Class, if applica			·						· -					
DUNS:	82-677-4775							_	Other Temperature Range	Requirement				
Proprietary Name (If Applicable) a	and Established Na	me: Famo	tidine Tablets USP, 20mg 10	00ct				I	(write in)					
Selling Unit NDC:	31722-017-10		Unit of Use NDC:				722017107		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Oral Solid - Table	t, Round Shaped, Li	ght Yellow, '11' and 'T'					I	Is this product to be shippe	d to customers on i	ice?		No	1
_									Is this product to be shippe				No	1
Active Ingredient(s):		Famotidine												
							b. Contact fo	r temperature excursion qu	estions:					
URL for Additional Product Inform								1	Name:		Soma Raju			
Address:		Ave (and) 800 Cente	ennial Ave, Suite 1		State:	Address 2:	00054	-	Number:		732-529-04			
City:	Piscataway Customer Service				State: Email:		08854	-	Group E-mail:		somaraju	@heterousa	a.com	
Key Contact: Phone Number:	1-866-827-3647	*			Fax:	732-562-8788	camberpharma.com	a Special re	gulations for product in any	ctatac?			No	7
Product Therapeutic Classification					ı ax.	732-302-0700		c. Special re	Special returns requiremen				No	-
Product Therapeutic Classification	on:								Special returns requiremen	is for this product?			INO	_
	ADDITI	ONAL PRODUCT I	NEORMATION			PRODUCT DESC	RIPTION INFORMATION	d Store prod	duct (unit of sale) upright?				No	7
	ADDITI	ONALT RODOUT II		Discoul Ohio Ool		T RODGOT DEGG	MI HOW IN CRIMATION	u. Store prot						4
The product is?			Is the Product	Direct-Ship Onl Neither	У		1000		Protect product (unit of s	ale) from light?			No	
a legend device? if yes, enter class #		No	Is the Product	Neither		Size:	1000ct	e. Shelf life:	Initial abolf life at larrach	if different).			24	Months Months
a product kit?		No	Orphan Drug Status				20mg		Initial shelf life at launch	ir ainerent):				Wonths
if yes, list NDCs of		INO	FDA Approval Status			Strength:	Zonig			ORDER INFOR	MATION			
component parts			- Bririppioral Glatao				Oral Solid - Tablet							
reverse numbered?		No				Dosage Form:			Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present						x Bottle		1 bottle of 1	000 tablets		
latex-free?		Yes				Product Shape:	Round		Box/Carton		(Write-in, e	.g. 1 Box of 1	0 Vials)	
preservative-free?		Yes				r routet snape.			Ampule					
correctional institution block?		No				Product Color:	Light Yellow		Glass		Minimum o	rder quantity	/?	Yes
opioid?		No							Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprint:	'11' and 'T'		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for	No	Lea Oble and advertise and a	and an the		•			Vial Liquid Multi Vial Powder Sql			Each	ich package	type?
hospital scanning? If Unit Dose, indicate NDC here:		NO	Is this product covered to Trade Agreements Act (		No				Vial Powder Sqi Vial Power Multi		24	Inner/Cartor	/Pook	
II Offic Dose, indicate NDC fiere.			Trade Agreements Act (	1747):	NO.				Other: Write In			Case	I/Fack	
			FOR GENERIC DRUG PR	ODUCTS				1	Other: Write III			Joase		
			TOR GENERIO DROGTT	000010										
					Au	thorized Generic *If A	authorized Generic, other		PI	HARMACY ORDER	R / BILL UNIT			
I. Orange Book Rating:	AB						ion fields are not applicable	Rec sell uni	t to customer?			nit to pharm	2011	
II. Generic Equivalent to What Bra		Pepcid						Tree. Sen um	t to customer i		KX billing u	Each	acy.	
ii. Generic Equivalent to What Bra	iliu:.	Горога						(Write-in, e.g	ı. 1 Vial)			Gram		
		DRUG SUPP	LY CHAIN SECURITY ACT	(DSCSA) INFORM	IATION				,			Milliliter		
				•								_		
Does supplier meet DSCSA defini	ition of manufactur	rer?	Yes		SLN:	0331722000000			ITE	II AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No											
If yes, select exemption:				0	SCP:			1		Dimens	ions (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:									Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No			iginal product purchase	ed	Item/Each:	0.3		2.25	4	0	1
Is product sold by manufacturer's			Yes	_	lirect from m						2.20		_ ·	' '
Has FDA granted waiver/exception		oduct?	No	<u>F</u>	Provide sour	ce manufacturer for rep	ackaged product	Box/Carton/i	Bundle/				0	
If yes, attach documentation fro	m FDA.							Inner Pack:						
		61	IN AND LUDGE PRODUCT I	NEODMATION				Case:	7.6	13.75	10	5.25	0	24
		GI	IN AND HIBCC PRODUCT I	NEORWATION				Pallet:		-				
Saleable Unit of Measure	c	Saleable Quantity	HIBCC		CTI	N-14	Unit of Use GTIN-14	Pallet:					0	
X Item/Each	3	1	TIIBUU			31722017107	Only Of USE Gillin-14			1		1	1	
Box/Carton/Bundle/Inner Pack					000				COST INFORMATION			WHOLESAL	ER USE ONL	
X Case		24			203	31722017101								
Pallet								Regular Cos	t		Vendor #:			
								Invoice Cost		\$100.00	Whsl. Code	#:		
								11			Fineline Co			
								As of date:			_			
								11						
ļ.								11						
			Attach copy of SAFETY D	ATA SHEET (SDS	) or non haza		ERT, LABEL AND PHOTO OF I	PRODUCT PACK						
*Please provide any additional inf	formation on page	2.				See new n 3 for Desi	gnated Drop Ship Only.		Signature:					



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

#### Version 2021

For Designated Drop Ship Only Products, Please Use Page 3

MA	TERIAL HAZ	ARD CLASSIFICATION and TRANSPORTATION				
Is this product (check all that apply): a. Cytotoxic?	SDS Hazard Classification					
<ul> <li>b. CA Prop. 65 Carcinogen or Reproductive Toxicant?</li> <li>Is the product a CA Prop 65 carcinogen?</li> <li>Is the product a CA Prop 65 reproductive toxicant?</li> <li>Does the product label bear a CA Prop 65 warning?</li> </ul>	X Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard					
c. Contact Hazard?      d. Does this product require special clean-up instructions?         (If yes, attach SDS with special instructions.)      e. Does the product contain DEHP?	Does the product have an Aerosol class? If yes, identify NFPA Storage Level:  NFPA Storage Level:					
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	No	Is the product a NIOSH hazardous drug? If yes, indicate which:	No			
c. DOT Hazard Class		Hazardous Waste Identification				
d. Packing Group e. Inhalation Hazard?	No	EPA Hazardous Waste Code:		Waste Characteristics		
Is this product regulated for shipment by IATA?	No		· · · · · · · · · · · · · · · · · · ·			
(if yes, answer a-e below and provide SDS) a. UN/Identification Number		REMS of	r REGISTRY RESTRICTIONS			
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	No	Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:	No			
Is the product restricted for air shipment? If so, indicate restriction:  Passenger Cargo Passenger & Cargo	No	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?)				
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?  (if yes, identify method below)  Limited Quantity  Consumer Commodity, ORM-D  Small Quantity (49 CFR 173.4)		REMS:  REMS Program Manager Name:  Supplier Manages REMS registry exclusively:  Wholesale distributor support:  Provider Name:  Site Enrollment Number assigned by Supplier:	No	Phone:  DEA #: NCPDP#: NPI #:		
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);		Comments				
SP#		Registry:	No			
ADD'L STORAGE INFORMATION		Registry Program Contact Name: Comments		Phone:		
Is the Product  Controlled Substance?  No Controlled Substance Code		R	ETURN INSTRUCTIONS			
Controlled by State(s)? ARCOS Reportable? Schedule No.  No Listed Chemical (List I or II) If yes, indicate which: Is it a scheduled listed chemical product?:	No	Contact tel. # if product received damaged:  Is product returnable for credit:	1-866-827-3647 Yes			
CLASS OF TRADE RESTRICTION:	Yes	URL/Link to returns policy:				
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	contact - customerservice@camberpharma.com					
Restricted to retail pharmacy only:  Restricted to hospital, clinics, and physician offices only:	No No	Special regulations or returns requirements for this product in certain states?				
Restricted from US territories? (explain in comments)  Comments:	If so, which states? Other requirements? Comments?					
M	SCELLANEC	OUS NOTES and/or Image of Product Barcode:				



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

#### Version 2021

#### FOR DESIGNATED DROP SHIP PRODUCT ONLY - if not a designated drop ship, do not complete.

Order Method for Designated Drop S	nip Product	Standard Order Receipt and Processing				
Purchase orders may be accepted by: a. EDI		Purchase order daily receipt cut off time by supplier Cut off time:				
b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	per:	Shipping lead time of PO:  Hours  Days  Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:				
Expedited Freight Charges or Other Designa	ed Drop Ship Fees:	Overnight and Priority Overnight PO Processing				
Expedited freight fees billed with each order:  Drop Ship service fee billed with each order:		Overnight receipt available:  PO Receipt cut off time:				
Drop Ship miscellaneous fees billed:  Comments:		Days of week overnight is available:  Monday Tuesday Wednesday Thursday Friday				
		Priority Overnight receipt available:				
Class of Trade Restriction		PO Receipt Cut off time:				
No restriction: Select YES if sold to retail pharmacy, hospitals, clinic Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	s and physician offices	Saturday Overnight receipt available:  PO Receipt Cut off time:  Phone: Fax: EDI:  Overnight Fees apply: Other fees apply:				
Other Data Information Required to F	rocess PO:	Return Instructions				
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty:		Contact # if product is received damaged:  Is product returnable for credit:  URL/Link to returns policy:  Special regulations or returns requirements for this product in certain states?  If so, which states? Other requirements? Comments?				
Miscellaneous Notes:						
		ADDITIONAL INFORMATION				
		Is product order for scheduled patient procedure? Is product order for restocking purposes?				