

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

Version 2021						Introduction Typ	e: New Item		x Final Version			Date:	11/2	7/2024
			PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOR	AGE REQUIF	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA					a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 214419				4419			Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)							
Medical Device Class, if applicable:														
DUNS:	11-856-3719								Other Temperature Range F	Requirement	Excursions p	ermitted to 1	5-30°C (59-8	86°F)
Proprietary Name (If Applicable) a	nd Established Na	ame: Ver	lafaxine Extended-Release Ta	blets 75 mg					(write in)	•				
Selling Unit NDC:	31722-124-90		Unit of Use NDC:		31722-124-90	UPC: 3	31722124904	1	Notes					
UDI			CVX Code:			MVX Code:								
Description: Venlafaxine Extended-Release Tablets 75 mg Is this product to be shipped to customers on ice? No									1					
									Is this product to be shipped				No	1
Active Ingredient(s): Venlafaxine hydrochloride, USP											-			-
								b. Contact for t	emperature excursion que	estions:				
URL for Additional Product Inform		www.camberpha	rma.com					4 1	Name:		Soma Raju			
Address:	800 Centennial Av					Address 2:			Number:		732-529-042			
City:	Piscataway				State:		Zip: 08854	Group E-mail:			somaraju@heterousa.com			
Key Contact:	1-866-827-3647	9			Email:	customerservice@c	amberpharma.com	c. Special regulations for product in any states?				NI.	1	
Phone Number:		0-1	and a construction of the last terms of the last	t (OND)	Fax:	732-562-8788					No	-		
Product Therapeutic Classification	1:	Selective serotonin	and norepinephrine reuptake inhib	tor (SNRI)					Special returns requirement	s for this product?			No	
	ADDITI	ONAL PROPUST	INFORMATION			BRODUOT DE	OODIDTION INFORMATION							7
	ADDITI	ONAL PRODUCT	INFORMATION			PRODUCT DE	SCRIPTION INFORMATION	d. Store produc	ct (unit of sale) upright?				No	
The product is?			Is the Product	Direct-Ship C	Only				Protect product (unit of sa	le) from light?			No	
a legend device?		No	Is the Product	Unit of Use		Size:	90 ct	e. Shelf life:					24	Months
if yes, enter class #			Orphan Drug Status					'	Initial shelf life at launch (if different):				Months
a product kit?		No	FD 4 4 1 0/-/			Strength:	75 mg		ORDER INFORMATION					
if yes, list NDCs of			FDA Approval Status				Futured ad selected films			ORDER INFORM	IATION			
component parts reverse numbered?		No				Dosage Form:	Extended-release, film coated tablet	II .	Unit of Sale		What is the	NDC calling	unit?	
co-licensed?		No	Allergens Present				coated tablet		x Bottle		1 Bottle of 90		uiiit:	
latex-free?		Yes	_				Round, biconvex		Box/Carton			g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Alc	ohol		Product Shape	rtouria, bicontox		Ampule		(**************************************	g. 1 Dox of 1	o vidio,	
correctional institution block?		No				Burnel Outer	White to off white		Glass		Minimum or	der quantity	?	Yes
opioid?		No				Product Color:			Tube					
Cannabinoid?		No	Country of Origin	USA		Product Imprin	'393' printed in black ink on one side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	nit dose for					Froduct imprin	and blank on the other side		Vial Liquid Multi		If Yes, how		ch package	type?
hospital scanning?			Is this product covered u						Vial Powder Sgl			Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (ΓAA)?	Yes				Vial Powder Multi			Inner/Carton	/Pack	
									Other: Write In			Case		
			FOR GENERIC DRUG PR	ODUCTS										
					Aut		f Authorized Generic, other			ARMACY ORDER	/ BILL UNIT			
	AB						ection fields are not applicable	Rec. sell unit to customer? Rx billing unit to pharmacy:						
II. Generic Equivalent to What Brand?: Venlafaxine Hydrochloride Extended-Release Tablets (Osmotica Pharmaceuti				a Pharmaceutica	al US LLC)		Each							
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION						(Write-in, e.g. 1 Vial)								
		DRUG SUP	PLY CHAIN SECURITY ACT	DSCSA) INFO	RMATION							Milliliter		
Does supplier meet DSCSA definit	ion of manufactur	ror2	Yes	_	GLN:	0331722498975			ITEN	I AND PACKING II	NEORMATION	J		
Is product exempt from DSCSA?	ion of manufactur	err	No	-	GLN.	0331722496973			11 = W	I AND I ACKING II	VI OKWATIOI	•		
*										D !	(110			
If yes, select exemption:					GCP:				Weight Lbs.		ons (US msm	•	Volume	Saleable # Pieces
Other exemption - Write in: Is product repackaged?			No		K			Item/Each:		Depth	Width	Height	(Cube)	Pieces
Is product repackaged? Is product sold by manufacturer's	evelucive dietribu	tor?	Yes	-	direct from mf	ginal product purcha	ised	item/Each:	0.12	2	2	3.75	15.00	1
Has FDA granted waiver/exception			No	-		e manufacturer for r	enackaged product	Box/Carton/Bu	ndle/					
If yes, attach documentation from		oudot.			Trovide Source	e manaradarer for f	сраскадса ргоциот	Inner Pack:	iluic,					
yoo, attaon accumentation from								Case:						
		G	TIN AND HIBCC PRODUCT I	NFORMATION				1	3.3	12.5	8.5	4	425.00	24
								Pallet:						
Saleable Unit of Measure	S	Saleable Quantity	HIBCC		GTIN	I-14	Unit of Use GTIN-14							
X Item/Each		1			0033	1722124904	00331722124904							
Box/Carton/Bundle/Inner Pack									COST INFORMATION		١	WHOLESALI	ER USE ONL	_Y:
X Case		24			1033	31722124901					l			
Pallet	1							Regular Cost			Vendor #:			
								Invoice Cost (V	VAC) (\$)	\$134.93	Whsl. Code			
								As of date:	3/1/2023		Fineline Cod	ie:		
								As of date:	3/ 1/2023		1			
	1													
1			Attach conv of SAFETY D	ATA SHEET (SE	S) or non hazar	d letter PACKAGE IN	SERT, LABEL AND PHOTO OF	PRODUCT PACKAG	SING and BARCODE		!			
*Please provide any additional info	ormation on page	2.	,doir dopy or or il ETT Dr	011221 (02	o, or non nazar		esignated Drop Ship Only.		Signature:					



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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL	HAZARD CLASSIFICATION and TRANSPORTATION					
Is this product (check all that apply): a. Cytotoxic? b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning?	SDS Hazard Classification x Organic Corrosive Oxidizer Inorganic Oxidizer 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? Is this product regulated for shipment by DOT?	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level: Is the product a NIOSH hazardous drug? No					
(if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class	If yes, indicate which: Hazardous Waste Identification					
d. Packing Group e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics					
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?	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? If Yes, is it managed with a pharmacy registry? Website URL:					
Is the product restricted for air shipment? If so, indicate restriction: Passenger Cargo Passenger & Cargo	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? No (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#: NCPDP#: NPI #:					
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments Registry: No					
	Registry Program Contact Name: Phone:					
ADD'L STORAGE INFORMATION	Comments					
Is the Product Controlled Substance? No Controlled Substance Code	RETURN 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 CLASS OF TRADE RESTRICTION:	Contact tel. # if product received damaged: 1-866-827-3647 Yes					
	URL/Link to returns policy:					
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Ye	contact - customerservice@camberpharma.com					
Restricted to retail pharmacy only:	Special regulations or returns requirements for this					
Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) No.	product in certain states? If so, which states? Other requirements? Comments?					
Comments:						
MISCELL	NEOUS NOTES and/or Image of Product Barcode:					



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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?