

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

Version 2021						Introduction Type:	Post Launch Change		x Final Version			Date:	11/20	0/2024
			PRODUCT INFORMA	ATION					SPECIAL HAN	IDLING AND STO	RAGE REQUI	REMENTS*		
Company Name:	Camber Pharmace	uticals, Inc.				Application:	ANDA	a. Temperatu	re - Indicate the USP temp	erature range for	this product.			
Application Number for NDA/AN	NDA/BLA (drug); PM	A/510(k)(med devi	ce):	212	2277				Temperature Range	Controlled Room		and 25 C (68	3° – 77° F)	
Medical Device Class, if applica								1	· -					
DUNS:	11-856-3719								Other Temperature Range	Requirement				
Proprietary Name (If Applicable)	and Established Nan	ne: Venlat	faxine Hydrochloride Extend	led-Release Cap	sules, USP 37.	5 mg (base)		I	(write in)					
Selling Unit NDC:	31722-002-30		Unit of Use NDC	:	31722-002-30		722002301		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Venlafaxine Hydrod	chloride Extended-R	Release Capsules, USP 37.5	mg (base)				Ţ	Is this product to be shippe	d to customers on	ice?		No	1
									Is this product to be shippe				No	1
Active Ingredient(s):		Venlafaxine hydroc	hloride, USP											
									r temperature excursion qu	estions:				
URL for Additional Product Inform		www.camberpharm	a.com						Name:		Soma Raju			
Address:	800 Centennial Ave	e, Suite 1			State:	Address 2:	00054		Number:		732-529-042			
City:	Piscataway Customer Service				State: Email:	customerservice@cam	08854		Group E-mail:		somaraju@h	eterousa.com		
Key Contact: Phone Number:	1-866-827-3647				Fax:	732-562-8788	iberpharma.com	a Special re	gulations for product in any	ctatos?			No	7
Product Therapeutic Classification		Saratonia and naran	inephrine reuptake inhibitor (	SVIDI)	I ax.	732-302-0700		c. Special re	Special returns requiremen				No	-
Product Therapeutic Classification	on:	Serotoriiri and norep	ineprime reuptake initibilor (	SINKI)					Special returns requiremen	is for this product?			INO	_
	ADDITIO	NAL PRODUCT IN	FORMATION			PRODUCT DESC	CRIPTION INFORMATION	I d Store prod	luct (unit of sale) upright?				No	7
	ADDITIO	NALT RODGOT IN		Discoul Ohio O	and it is	T RODGOT DEGG	ANI TISH IN SKIIIATISH	u. Store prot						4
The product is?	ı	NI.	Is the Product	Direct-Ship O Unit of Use	inly		20 -1	- 01-14-14	Protect product (unit of s	ale) from light?			No	
a legend device? if yes, enter class #		No	Is the Product	Offic of Ose		Size:	30 ct	e. Shelf life:	luitiel abalf life at launch	if different).			24	Months Months
a product kit?		No	Orphan Drug Status				37.5 mg		Initial shelf life at launch	ir airierent):				Wonths
if yes, list NDCs of		INO	FDA Approval Status			Strength:	57.5 mg			ORDER INFOR	MATION			
component parts			. Ditirippi oral otatao				Hard gelatin capsule							
reverse numbered?		No				Dosage Form:	3		Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present						x Bottle		1 Bottle of 3	0 Capsules		
latex-free?		Yes	Gluton Dvo	Alcohol, Sugar		Product Shape:	Capsule		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Oluten, Dye,	Alconol, Gugui		i roduct onapc.			Ampule					
correctional institution block?		No				Product Color:	Grey opaque cap & white		Glass		Minimum o	rder quantity	y?	Yes
opioid?		No					opaque body		Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprint:	Imprinted with 'V' on cap & '9' on body printed in black color		Vial Liquid Sgl					
If Unit Dose, is item bar coded to	unit dose for		In this was done a service of	and an the		•	on body printed in black color		Vial Liquid Multi				ich package	type?
hospital scanning?			Is this product covered Trade Agreements Act		No				Vial Powder Sgl Vial Powder Multi		24	Each Inner/Cartor	-/Dools	
If Unit Dose, indicate NDC here:	Į.		Trade Agreements Act	(IAA)!	INO				Other: Write In			Case	1/Pack	
			FOR GENERIC DRUG PR	PODUCTS				J	Other: Write III			Ousc		
			TOR GENERIC DROG FI	CODOCIO										
					Au	thorized Generic *If A	Authorized Generic, other		PI	IARMACY ORDER	R / BILL UNIT			
I Oronno Book Botinos	AB				,,,,		tion fields are not applicable	Pac sall uni	to customer?					
I. Orange Book Rating: II. Generic Equivalent to What Bra		Effexor XR					•••	ixec. sen um	to customer:	1	Rx billing u	Each	acy:	
II. Generic Equivalent to What Bra	aliur.	LIIEXUI XIX						(Write-in, e.g	1 Vial)			Gram		
		DRUG SUPPI	LY CHAIN SECURITY ACT	(DSCSA) INFOR	MATION			(**************************************				Milliliter		
				, ,										
Does supplier meet DSCSA defin	ition of manufacture	er?	Yes		GLN:	0331722498975			ITEI	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No											
If yes, select exemption:	Ī				GCP:			1		Dimens	ions (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:									Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was or	iginal product purchase	ed	Item/Each:	0.08	1.5	1.5	3	6.75	1
Is product sold by manufacturer's			Yes		direct from m					1.5	1.0	,	0.73	'
Has FDA granted waiver/exception		duct?	No		Provide source	ce manufacturer for rep	ackaged product	Box/Carton/I	Bundle/					
If yes, attach documentation fro	om FDA.							Inner Pack:						
		CT	IN AND HIBCC PRODUCT	NEODMATION				Case:	2.05	9.5	6.75	4	256.50	24
		GII	IN AND RIBCC PRODUCT	INFORMATION				Pallet:			-			
Saleable Unit of Measure	Ç.,	leable Quantity	HIBCC		CTII	N-14	Unit of Use GTIN-14	Pallet:						
X Item/Each	Sa	1	ПІВСС			31722002301	00331722002301							
Box/Carton/Bundle/Inner Pack	ŀ				003	3 <u>220020</u> 01	00001122002001		COST INFORMATION			WHOLESAL	ER USE ONL	LY:
X Case		24			303	31722002302								
Pallet					1   200			Regular Cos	t		Vendor #:			
								Invoice Cost		\$6.80	Whsl. Code	#:		
											Fineline Co	de:		
								As of date:	12/1/2024					
											-			
									-					
*Please provide any additional in			Attach copy of SAFETY D	ATA SHEET (SD	S) or non haza		ERT, LABEL AND PHOTO OF F	PRODUCT PACK	AGING and BARCODE.  Signature:					



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

#### Version 2021

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 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?  Is this product regulated for shipment by DOT?	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					
<del></del>	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:					



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

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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?