

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

Version 2021					Introduction 1	Туре:	New Item	0000	Final Version			Date:	11/28/2	2023	
PRODUCT INFORMATION						SPECIAL HANDLING AND STORAGE REQUIREMENTS*									
Company Name: Camber Pharmaceuticals, Inc.				Applica	tion:	ANDA	a. Temperature – Indicate the USP temperature range for this product.								
	/ANDA/BLA (drug); PMA/510(k)(med device): 214571					Temperature Range Controlled Room – between 20 and 25 C (68° – 77°					° – 77° F)				
Medical Device Class, if applicable:															
	11-856-3719							Other Te	emperature Range F	Requirement	Excursions p	ermitted to 1	5° to 30° C (59	9° – 86° F)	
Proprietary Name (If Applicable) and		Varenicline Tablets 0.5 mg &							rite in)						
	31722-690-31	Unit of Use N	DC:	31722-690-31	UPC:	3317226	590317	Notes							
UDI		CVX Code:			MVX Code:										
Description:	Varenicline Tablets 0.5 mg and 1 m	ng with "H" on one side and "\/23" on the other	side 1 mg - Vellow deb	occed with "H" on o	ne side and "\/?/" on the	other side			oduct to be shipped				No		
**NOTE - 0.5 mg - Pink, debossed with "H" on one side and "V23" on the other side. 1 mg - Yellow, debossed with "H" on one side and "V24" on the other side. Is this product to be shipped to customers on dry ice? No Active Ingredient(s): Varenicline Tartrate															
Active ingredient(s): valencine raturate b. Contact for temperature excursion questions:															
URL for Additional Product Information: www.camberoharma.com								Name: Soma Raju							
Address:	1031 Centennial Ave (and)	031 Centennial Ave (and) 800 Centennial Ave, Suite 1			Address 2:				Number:			732-529-0423			
City:	Piscataway			State:	NJ	Zip:	08854	Group E-mail: somaraju@heter			eterousa.com	erousa.com			
-	Customer Service			Email:		omerservice@camberpharma.com									
Phone Number:	1-866-827-3647			Fax:	732-562-8788			c. Special regulations for product in any states? No							
Product Therapeutic Classification	cation: Nicotinic receptor partial agonist							Special returns requirements for this product? No							
	ADDITIONAL PRODUCT INFORMATION PRODUCT DESCRIPTION INFORMATION d. Store product (unit of sale) upright? No														
	ADDITIONAL PR		Distant Citi	0	PRODUCT	DESCRIP	HON INFORMATION		,				No		
The product is?	N	Is the Product	Direct-Ship Unit of Use	Unly					product (unit of sa	le) from light?			No	Manutha	
a legend device? if yes, enter class #	No	Is the Product Orphan Drug Statu			Size:	5	i3 ct	e. Shelf life:	nelf life at launch (i	f different).			24	Months Months	
a product kit?	No	Orphan Drug Statu				0.	.5 mg & 1 mg	initial Si	ien nie at launch (i	i unerentj.				wonths	
if yes, list NDCs of		FDA Approval Stat	IS		Strength:					ORDER INFORM	IATION				
component parts					Dosage Form	m. Fi	ilm coated tablets								
reverse numbered?	No				Dosageron			Unit of S			What is the I		unit?		
co-licensed?	No	Allergens Present				_			Bottle		1 Carton of 5				
latex-free?	Yes		Alcohol		Product Sha	ape: C	Capsular, biconvex	X	Box/Carton		(Write-in, e.g	g. 1 Box of 10	Vials)		
preservative-free? correctional institution block?	Yes	_				**	*See Note in Descripton		Ampule Glass		Minimum or	dor quantitu	,	Yes	
opioid?	No	_			Product Col	or: **	*		Tube		Minimum or	der quantity	ſ	res	
Cannabinoid?	No	Country of Origin	India			**	*See Note in Descripton		Vial Liquid Sgl						
If Unit Dose, is item bar coded to un					Product Imp	wint:	*	Vial Liquid Multi If Yes, how many of which package type?						/pe?	
hospital scanning?		Is this product cove						Vial Powder Sql 48 Each							
If Unit Dose, indicate NDC here:		Trade Agreements A	ct (TAA)?	No				Vial Power Multi Inner/Carton/Pack							
									Other: Write In			Case			
		FOR GENERIC DRUC	PRODUCTS					_							
				Au	thorized Generic	*If Autho	orized Generic, other	PHARMACY ORDER / BILL UNIT							
I. Orange Book Rating:	AB						fields are not applicable	Rec. sell unit to custo				hit to pharma	cv.		
I. Generic Equivalent to What Brand?: Chantix							Rec. sell unit to customer? Rx billing unit to pharmacy:								
· · · · · · · · · · · · · · · · · · ·							(Write-in, e.g. 1 Vial) Gram								
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION Milliliter															
Does supplier meet DSCSA definit	ion of monufactures?	Yes		GLN:	0331722000000				ITEM	AND PACKING I		1			
Is product exempt from DSCSA?	ion of manufacturer?	No		GLN:	0331722000000					AND PACKING I	NFORMATION				
				GCP:						Dimorra	ons (US msm	te)	Value	Calaah I. #	
If yes, select exemption: Other exemption - Write in:				GCP:					Weight Lbs.	Dimensi	Width	Height	Volume (Cube)	Saleable # Pieces	
Is product repackaged?		No		If yes, was o	riginal product pur	chased		Item/Each:	0.07				· /		
Is product sold by manufacturer's	exclusive distributor?	Yes		direct from m					0.27	7.4	1.3	4	38.48	1	
Has FDA granted waiver/exception		No		Provide sour	ce manufacturer fo	or repacka	aged product	Box/Carton/Bundle/							
If yes, attach documentation from	n FDA.							Inner Pack:							
		GTIN AND HIBCC PRODU						Case:	13.6	15.25	11.5	12.5	2,192.19	48	
		GTIN AND HIBCC PRODU	TINFORMATION					Pallet:							
Saleable Unit of Measure	Saleable Q	uantity HIBCC		GTI	N-14		Unit of Use GTIN-14								
X Item/Each	1				31722690317		00331722690317		1	1					
Box/Carton/Bundle/Inner Pack	n/Bundle/Inner Pack							COS	T INFORMATION		١	WHOLESALE	R USE ONLY	/:	
X Case	48			203	31722690311	_									
Pallet						-		Regular Cost			Vendor #:				
						-		Invoice Cost (WAC) (\$		\$207.78	Whsl. Code				
				-		-		As of date:	1/11/2024		Fineline Coo	<i>i</i> c.			
						-		, is of date.			1				
L		Attach copy of SAFET	/ DATA SHEET (S	DS) or non haza	rd letter, PACKAGE	INSERT,	LABEL AND PHOTO OF P	RODUCT PACKAGING an	d BARCODE.		•				
*Please provide any additional info	ormation on page 2.		(-	,			ted Drop Ship Only.	Signatu							

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

Version 2021 For Des	signated Drop Ship Only Products, Please Use Page 3						
MATERIA	L HAZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply): a. Cytotoxic? N b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? N Is the product a CA Prop 65 reproductive toxicant? N Does the product label bear a CA Prop 65 warning? 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? N	x Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Image: Contact Hazard						
e. Does the product contain DET if it is product regulated for shipment by DOT? 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?	o 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?	Is there a REMS on this product? No 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	o Med Guide Required No Limited Distribution Requirement Image: Comments / Details: (For example, iPledge program?) Image: Comment of the second secon						
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	REMS: No REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: Provider Name: Provider Name: DEA #: Site Enrollment Number assigned NCPDP#: by Supplier: NPI #:						
Special Provision (listed in Column 7 of 49 CFR 172.101);	Parister No.						
SP#ADD'L STORAGE INFORMATION	Registry: No Registry Program Contact Name: Phone: Comments						
Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS						
Controlled by State(s)? ARCOS Reportable? Schedule No. No CLASS OF TRADE RESTRICTION: No No CLASS OF TRADE RESTRICTION:	Contact tel. # if product received damaged: 1-866-827-3647						
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	contact - customerservice@camberpharma.com						
No resultation: seed Yes if source retail pharmacy, nospitals, cuinces and physician offices If the second seed Yes if source retail pharmacy, nospitals, cuinces and physician offices Restricted to retail pharmacy only: N Restricted to hospital, clinics, and physician offices only: N Restricted from US territories? (explain in comments) N Comments: Image: Comment in the second se	O Special regulations or returns requirements for this o product in certain states?						
MISCEL	ANEOUS NOTES and/or Image of Product Barcode:						



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Version 2021 FOR DESIGNATED DROP SHIP PRODUCT ONLY - if	not a designated drop ship, do not complete.
Order Method for Designated Drop Ship Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name: Phone:	Purchase order daily receipt cut off time by supplier Cut off time: 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 Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order: Drop Ship miscellaneous fees billed: Comments:	Overnight receipt available: Image: Comparison of the co
Class of Trade Restriction:	PO Receipt Cut off time:
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	Saturday Overnight receipt available: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to Process PO:	Return Instructions
Patient Procedure Date:	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?