

# **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	TION					SPECIAL HAN	DLING AND STO	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA						a. Temperature – Indicate the USP temperature range for this product.								
Application Number for NDA/AN	NDA/BLA (drug); PN	/IA/510(k)(med devi	ce):	20	4389				Temperature Range	Controlled Room		and 25 C (68	3° – 77° F)	
Medical Device Class, if applica			·						· -					
DUNS:	11-856-3719								Other Temperature Range I	Requirement				
Proprietary Name (If Applicable)	and Established Na	me: Mema	antine Hydrochloride Tablets						(write in)					
Selling Unit NDC:	31722-807-60		Unit of Use NDC		31722-807-60		722807609		Notes					
UDI			CVX Code:			MVX Code:								
Description:	Memantine Hydro	chloride Tablets, US	iP 5 mg						Is this product to be shipped	d to customers on	ice?		No	1
									Is this product to be shipped				No	1
Active Ingredient(s):		Memantine hydrocl	hloride, USP											
								b. Contact fo	r temperature excursion qu	estions:				
URL for Additional Product Inform		www.camberpharm	na.com						Name:		Soma Raju			
Address:	800 Centennial Av	ve, Suite 1			State:	Address 2: NJ Zin	00054		Number:		732-529-042			
City:	Piscataway Customer Service				State: Email:	customerservice@cam	08854		Group E-mail:		somaraju@f	neterousa.com	<u>n</u>	
Key Contact: Phone Number:	1-866-827-3647	!			Fax:	732-562-8788	iberpharma.com	a Special rea	gulations for product in any	ctatos?			No	1
Product Therapeutic Classification		N-methyl-D-separts	ate (NMDA) receptor antago	niet	I ax.	732-302-0700		c. Special reg	Special returns requirement				No	-
Product Therapeutic Classification	on:	IN-IIIetilyi-D-aspaita	ate (INIVIDA) receptor antago	1151					Special returns requirement	s for this product?			INO	_
	ADDITIO	ONAL PRODUCT IN	JEORMATION			PRODUCT DESC	RIPTION INFORMATION	d Store prod	uct (unit of sale) upright?				No	1
	ADDITI	ONALT RODUCT III		Discoul Object	2-1-	TRODUCT DECC	IKII TION IN OKMATION	u. Store prou						1
The product is?		NI.	Is the Product	Direct-Ship ( Unit of Use	only		00 -1	. 01-1/17	Protect product (unit of sa	ile) from light?			No	
a legend device? if yes, enter class #		No	Is the Product Orphan Drug Status	Offic of Ose		Size:	60 ct	e. Shelf life:	Initial shelf life at launch (	if different).			24	Months Months
a product kit?		No	Orphan Drug Status				5 mg		initial shell life at launch (	ir different):				Wonths
if yes, list NDCs of		140	FDA Approval Status			Strength:	5 mg			ORDER INFOR	MATION			
component parts			- Dririppioral Glatag				Film coated 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 6	0 Tablets		
latex-free?		Yes	Dairy, Lactose, Cas	ain Dve Corn	Alcohol	Product Shape:	Modified capsule,		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Dan y, Luciosc, Ous	om, byc, com,	Alcohol	i roddot onapc.	biconvex		Ampule					
correctional institution block?	•	No				Product Color:	Tan		Glass		Minimum o	der quantity	/?	Yes
opioid?		No							Tube					
Cannabinoid?		No	Country of Origin	India		Product Imprint:	Debossed with 'J' on one side & '47' on the other side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to	unit dose for		Le Oble and death account	and an the			side & 47 on the other side		Vial Liquid Multi				ich package t	type?
hospital scanning?			Is this product covered Trade Agreements Act		No				Vial Powder Sgl Vial Powder Multi		24	Each	/Deels	
If Unit Dose, indicate NDC here:			Trade Agreements Act	IAA)!	INO				Other: Write In			Inner/Cartor Case	I/Pack	
			FOR GENERIC DRUG PR	ODUCTS					Other: Write in			Ousc		
			TOR GENERIC DROG FI	000013										
					Aut	horized Generic *If A	uthorized Generic, other		PH	ARMACY ORDER	R / BILL UNIT			
I Oronno Book Botings	AB				/		ion fields are not applicable	Pac sall unit	to customer?					
I. Orange Book Rating: II. Generic Equivalent to What Bra		Namenda						Nec. sen unit	to customer:	1	Rx billing u	Each	acy:	
II. Generic Equivalent to What Bra	anur.	Ivamenda						(Write-in, e.g	1 Vial)	1		Gram		
		DRUG SUPP	LY CHAIN SECURITY ACT	(DSCSA) INFO	RMATION			(**************************************				Milliliter		
				,										
Does supplier meet DSCSA defin	nition of manufactur	er?	Yes		GLN:	0331722498975			ITEN	I AND PACKING I	INFORMATIO	4		
Is product exempt from DSCSA?	•		No						·					
If yes, select exemption:					GCP:					Dimens	ions (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:								1	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If yes, was or	iginal product purchase	ed	Item/Each:	0.05	1.5	1.5	2.52	5.67	1
Is product sold by manufacturer's	's exclusive distribu	itor?	Yes		direct from m	fr?			0.00	1.5	1.5	2.02	3.07	' '
Has FDA granted waiver/exception		oduct?	No		Provide source	e manufacturer for repart	ackaged product	Box/Carton/E	Bundle/					
If yes, attach documentation fro	om FDA.							Inner Pack:						
		0.7	IN AND HIDDO PRODUCT	NEODMATION				Case:	1.9	9.5	6.9	4	262.20	24
		GI	IN AND HIBCC PRODUCT	NFORMATION				Dellet						
Saleable Unit of Measure	c	aleable Quantity	HIBCC		GTIN	J-14	Unit of Use GTIN-14	Pallet:						
X Item/Each	5	aleable Quantity	ПІВСС			N-14 31722807609	00331722807609							
Box/Carton/Bundle/Inner Pack		•			3030	22307000			COST INFORMATION			WHOLESAL	ER USE ONL	Y:
and an		24			2033	31722807603								
X Case								Regular Cost			Vendor #:			
X Case Pallet								Invoice Cost		\$8.60	Whsl. Code	#:		
								1.1						
											Fineline Co	de:		
								As of date:	12/1/2024		Fineline Co	de:		
								As of date:	12/1/2024		Fineline Co	de:		
											Fineline Co	de:		
			Attach copy of SAFETY D	ATA SHEET (SD	OS) or non hazar		ERT, LABEL AND PHOTO OF P				Fineline Co	de:		



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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 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:					



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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?