

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

Version 2021						Introduction ¹	Type:	Post Launch Change] [x Final Version			Date:	8/26/	2024
			PRODUCT INFORMAT	TION						SPECIAL HAND	LING AND STOR	AGE REQUIF	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc.					Applica	Application: ANDA			a. Temperature – Indicate the USP temperature range for this product.						
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): Ognocolumnaria (assertion of the perturbation of the pert															
Medical Device Class, if applicable:															
DUNS:	11-856-3719								1	Other Temperature Range F	Requirement				
Proprietary Name (If Applicable) a		me: Lithiu	um Carbonate Capsules, USP	300 mg					1	(write in)					
Selling Unit NDC:	31722-545-01		Unit of Use NDC:	J		UPC:	33172254	5013		Notes					
UDI			CVX Code:			MVX Code:									
Description:	Lithium Carbonate	Cansules USP 30	00 ma	-					i	Is this product to be shipped	I to customers on i	ne?		No	
Is this product to be shipped to											No				
Active Ingredient(s):		Lithium carbonate	e, USP									,			
b. Contact for temperature excursion questions:															
URL for Additional Product Information: www.camberpharma.com							Name: Soma Raju								
Address:	800 Centennial Av	ennial Ave, Suite 1				Address 2:			Number: 732-529-0423						
City:	Piscataway	State:			NJ	Zip: 08		Group E-mail: somaraju@heterousa.com					<u>m</u>		
Key Contact:	Customer Service	Email:				customerservice@camberpharma.com									
Phone Number:	1-866-827-3647		Fax			732-562-8788				lations for product in any				No	
Product Therapeutic Classificatio	n:	Antimanic mood s	stabilizing agent							Special returns requirements	s for this product?			No	
	ADDITIC	NAL PRODUCT I	NFORMATION			PRODUCT	DESCRIPTI	ON INFORMATION	d. Store produc	ct (unit of sale) upright?				No	
The product is?			Is the Product	Direct-Ship C	nly					Protect product (unit of sa	le) from light?			No	
a legend device?		No	Is the Product	Neither		Size:	100) ct	e. Shelf life:					24 Months	Months
if yes, enter class #			Orphan Drug Status			0.20.				Initial shelf life at launch (i	f different):				Months
a product kit?		No				Strength:	300) mg							
if yes, list NDCs of	FDA Approval Status					Hard gelatin capsule				ORDER INFORM	IATION				
component parts						Dosage For	Dosage Form:			U-1-1-4 O-1-		What is the	NDC aallina		
reverse numbered? co-licensed?		No	Allermana Dracent							Unit of Sale x Bottle		1 Bottle of 10		unitr	
latex-free?	No Allergens Present					Capsule									
preservative-free?		Yes	Alcohol, Ani	mal Products		Product Sha	ape:	osule	Box/Carton (Write-in, e.g. 1 Box of 10 V Ampule				o viais)		
correctional institution block?		No					Pin	k body and pink cap		Glass		Minimum or	der quantity	12	Yes
opioid?		No				Product Col	lor:	k body and pink oup		Tube			uoi quantiti	,.	103
Cannabinoid?		No	Country of Origin	India			. Imr	printed with '98' on		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u			,			Product Imp		ly and 'H' on cap		Vial Liquid Multi		If Yes, how	many of whi	ich package	type?
hospital scanning?			Is this product covered u	nder the						Vial Powder Sgl			Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (TAA)?	No							Inner/Carton	/Pack		
										Other: Write In			Case		
			FOR GENERIC DRUG PRO	ODUCTS											
												-			
					Au	thorized Generic		zed Generic, other		PH.	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB					section fields are not applicable			Rec. sell unit to customer?			Rx billing unit to pharmacy:			
II. Generic Equivalent to What Bra		Lithium Carbonate	e (Roxane)							Each					
-		-							(Write-in, e.g. 1 Vial)			Gram			
		DRUG SUPPI	LY CHAIN SECURITY ACT (DSCSA) INFOR	MATION								Milliliter		
Does supplier meet DSCSA defini	ition of manufactur	rer?	Yes		GLN:	0331722498975				ITEM	AND PACKING IN	IFORMATION			
Is product exempt from DSCSA?			No												
If yes, select exemption:					GCP:					Weight Lbs.	Dimensi	ons (US msm	its.)	Volume	Saleable #
Other exemption - Write in:										Troigin LDS.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No			iginal product			Item/Each:	0.17	2.2	2.2	4	19.36	1
Is product sold by manufacturer's				1		rect from mfr?						-			
			Yes	_											
Has FDA granted waiver/exceptio	n/exemption for pr		Yes No		Provide sour	ce manufacturer f	for repacka	ged product	Box/Carton/Bu	ndle/					
	n/exemption for pr				Provide sour	ce manufacturer f	for repacka	ged product	Inner Pack:	ndle/				653.13	24
Has FDA granted waiver/exceptio	n/exemption for pr	oduct?	No		Provide sour	ce manufacturer f	for repacka	ged product		4.65	13.75	9.5	5	653.13	2-7
Has FDA granted waiver/exceptio	n/exemption for pr	oduct?			Provide sour	ce manufacturer f	for repacka	ged product	Inner Pack: Case:		13.75	9.5	5	003.13	
Has FDA granted waiver/exceptio If yes, attach documentation fro	on/exemption for pr om FDA.	oduct?	No IN AND HIBCC PRODUCT IN						Inner Pack:		13.75	9.5	5	653.13	
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure	on/exemption for pr om FDA.	oduct?	No		GTII	N-14		nit of Use GTIN-14	Inner Pack: Case:		13.75	9.5	5	653.13	2-7
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure x tem/Each	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN		GTII				Inner Pack: Case:		13.75			ER USE ONL	
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X tem/Each Box/Carton/Bundle/Inner Pack	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN		GTII 0033	N-14 31722545013			Inner Pack: Case:	4.65	13.75				
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure x tem/Each	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN		GTII 0033	N-14			Inner Pack: Case:	4.65	13.75				
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X Item/Each Box/Carton/Bundle/Inner Pack X Case	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN		GTI 003:	N-14 31722545013			Inner Pack: Case: Pallet:	4.65 COST INFORMATION		V	VHOLESALI		
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X Item/Each Box/Carton/Bundle/Inner Pack X Case	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN		GTI 003:	N-14 31722545013			Inner Pack: Case: Pallet: Regular Cost	4.65 COST INFORMATION VAC) (\$)		Vendor #:	VHOLESALI		
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X Item/Each Box/Carton/Bundle/Inner Pack X Case	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN		GTI 003:	N-14 31722545013			Inner Pack: Case: Pallet: Regular Cost	4.65 COST INFORMATION		Vendor #: Whsl. Code	VHOLESALI		
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X Item/Each Box/Carton/Bundle/Inner Pack X Case	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN		GTI 003:	N-14 31722545013			Inner Pack: Case: Pallet: Regular Cost Invoice Cost (V	4.65 COST INFORMATION VAC) (\$)		Vendor #: Whsl. Code	VHOLESALI		
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X Item/Each Box/Carton/Bundle/Inner Pack X Case	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN HIBCC	IFORMATION	GTII 0033 2033	N-14 31722545013 31722545017	U	nit of Use GTIN-14	Inner Pack: Case: Pallet: Regular Cost Invoice Cost (V	4.65 COST INFORMATION VAC) (\$) 9/28/2010		Vendor #: Whsl. Code	VHOLESALI		
Has FDA granted waiver/exceptio If yes, attach documentation fro Saleable Unit of Measure X Item/Each Box/Carton/Bundle/Inner Pack X Case	on/exemption for pr om FDA.	GTI aleable Quantity	No IN AND HIBCC PRODUCT IN	IFORMATION	GTII 0033 2033	N-14 31722545013 31722545017	U	nit of Use GTIN-14	Inner Pack: Case: Pallet: Regular Cost Invoice Cost (V	4.65 COST INFORMATION VAC) (\$) 9/28/2010		Vendor #: Whsl. Code	VHOLESALI		



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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HA	ZARD 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? No	SDS Hazard Classification x Organic Corrosive					
Is the product a CA Prop 65 reproductive toxicant? Does the product label bear a CA Prop 65 warning? C. Contact Hazard? No	Inorganic Oxidizer Contact Hazard					
d. Does this product require special clean-up instructions? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No	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	Is the product a NIOSH hazardous drug? If yes, indicate which:					
c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics					
Is this product regulated for shipment by IATA?						
(if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS					
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? 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:					
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 Controlled by State(s)? No Listed Chemical (List I or II) No	RETURN INSTRUCTIONS					
ARCOS Reportable? Schedule No. If yes, indicate which: Is it a scheduled listed chemical product?: No	Contact tel. # if product received damaged: Is product returnable for credit: Yes					
CLASS OF TRADE RESTRICTION:	URL/Link to returns policy:					
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Restricted to retail pharmacy only: No	contact - customerservice@camberpharma.com					
Restricted to retail pharmacy only. Restricted to hospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No	Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?					
Comments:	1 1 30, Willow States. Other requirements: Comments:					
MISCELLANE	OUS 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 Ship Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI b. Autofax Fax Number:	Purchase order daily receipt cut off time by supplier Cut off time:
c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number:	Shipping lead time of PO: Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:
Contracted 3PL company / contact #: Name: Phone:	
Expedited Freight Charges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order:	Overnight receipt available:
Drop Ship service fee billed with each order:	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, 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: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to Process 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?