

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

Version 2024						Introduction 1	Type: New li	tem		x Final Version			Date:	4/7/2	2025
			PRODUCT INFORMAT	TION						SPECIAL HAN	IDLING AND STOP	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc.				Application: ANDA		a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/AND	DA/BLA; PMA/510(k):	218272	2			NDA 505(b) Type	NOT APPLICA	ABLE		mperature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicab									í I						
	11-856-3719								Otl	ner Temperature Range	Requirement	Excursions	permitted to 1	5° to 30°C (5	9° to 86°F)
Proprietary Name (If Applicable) and		Chlorpr	omazine Hydrochloride Injec	ction, USP 25 m	ng/mL (Single-D					(write in)					
Selling Unit NDC:	31722-366-32		Unit of Use NDC:			UPC: MVX Code:	331722366328		No	tes					
U 2.			CVX Code:			WVX Code.									
Description:	Chlorpromazine Hydroch	loride Injection	, USP 25 mg/mL (Single-Do	se Vials)						his product to be shippe				No	
Active Ingredient(s): Chlorpromazine hydrochloride, USP							IST	his product to be shippe	d to customers on c	iry ice?		No			
Active ingreutency. On opportable ingreutency of						b. Contact for ten	perature excursion qu	estions:							
URL for Additional Product Inform	ation: www.	camberpharma	.com							me:		Soma Raju			
Address:	800 Centennial Ave, Suite 1			Address 2:			Nu	732-529-0423							
	Piscataway				NJ Zip: 08854			Group E-mail: somaraju@heterousa.com							
Key Contact:		Customer Service Email:			customerservice@camberpharma.com			c. Special regulations for product in any states? No							
Phone Number:	1-866-827-3647				Fax:	732-562-8788								No	
Product Therapeutic Classification	Anup	sychotic drug]				Sp	ecial returns requiremen	ts for this product?			No	
	ADDITIONAL	PRODUCT INF	ORMATION			PRODUCT	DESCRIPTION INFORM	ATION	d Store product	(unit of sale) upright?				No	1
The product is?			Is the Product	Direct-Ship O	iniu	111020011				otect product (unit of s	ala) from links?			No	
a legend device?	No		Is the Product	Unit Dose	any		25 x 1 mL singl	e-dose	e. Shelf life:	Steet product (unit of S	ale) from light?			24	Months
if yes, enter class #			Orphan Drug Status	Crine Docto		Size:	vials	0 0000		tial shelf life at launch	(if different):			27	Months
a product kit?	No					Strength:	25 mg/mL per single-dose								
if yes, list NDCs of			FDA Approval Status			Strength.	vial				ORDER INFORM	IATION			
component parts						Dosage Form	m: Sterile, clear sol	ution							
reverse numbered? co-licensed?	No		Allermone Dresent			•			Un	it of Sale			NDC selling		le.
co-licensed? latex-free?	No Yes		Allergens Present				N/A			x Box/Carton			25 x 1 mL Sin .g. 1 Box of 1		IS
preservative-free?	Yes					Product Sha	ape:			Ampule		(11111111111111111111111111111111111111	.g. i box oi ii	5 viais)	
correctional institution block?	No					Product Col	Colorless			x Glass		Minimum o	rder quantity	?	Yes
opioid?	No					Product Col	or:			Tube					
Cannabinoid?	No		Country of Origin	India		Product Imp	n/A			x Vial Liquid Sgl					
If Unit Dose, is item bar coded to un										Vial Liquid Multi			many of whi	ch package	type?
hospital scanning? If Unit Dose, indicate NDC here:	Yes	2-366-31	Is this product covered un Trade Agreements Act (T		No					Vial Powder Sgl Vial Powder Multi		1	Each Inner/Carton	Pack	
in onit bose, indicate type here.	5172	2-300-31	Thate Agreements Act (1	700	NO					Other: Write In			Case	/I dok	
			FOR GENERIC DRUG PRO	ODUCTS											
					Au	thorized Generic	*If Authorized Generic			Pł	HARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AP						section fields are not a	applicable	Rec. sell unit to customer?			Rx billing unit to pharmacy: Each			
II. Generic Equivalent to What Bran	nd?: Chlor	promazine Hyd	rochloride Injection, USP RI	D of Hikma Ph	armaceuticals I	USA Inc.									
									(Write-in, e.g. 1 Vial) Gram						
		DRUG SUPPL	Y CHAIN SECURITY ACT (I	DSCSA) INFOR	MATION				HCPCS J-Code:	J3230			Milliliter		
Does supplier meet DSCSA definit	ion of manufacturer?		Yes	Т	GLN:	0331722498975					M AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?		1	No	-											
If yes, select exemption:				_	GCP:						Dimensi	ons (US msr	nts.)	Volume	Saleable #
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No			iginal product pur	chased		Item/Each:	0.43	3.38	3.38	2.25	25.70	1
Is product sold by manufacturer's			Yes	-	direct from m						0.00	0.00	2.20	20.10	
Has FDA granted waiver/exception		?	No		Provide source	ce manufacturer fo	or repackaged product		Box/Carton/Bund	le/					
If yes, attach documentation from	II DA.								Inner Pack: Case:						
		GTIN	AND HIBCC PRODUCT IN	FORMATION					ouse.	8.65	11	11	5.5	665.5	18
									Pallet:						
Saleable Unit of Measure	RFID tag(Y/N) Salea		HIBCC		GTI	N-14	Unit of Use G	TIN-14							
	Quan	tity			000	31722366328									
X Item/Each Box/Carton/Bundle/Inner Pack	N	1			003	31722366328				COST INFORMATION			WHOLESAL	ER LISE ONI	v .
X Case	N	18			203	31722366322							MINOLEOAL		
Pallet									Regular Cost			Vendor #:			
									Invoice Cost (WA	C) (\$)	\$624.00	Whsl. Code	#:		
							_					Fineline Co	de:		
							_		As of date:	4/1/2025		-			
┝╋			Attach copy of SAFETY DA		S) or non haza							1			
*Please provide any additional info	ormation on page 2.		, and on oupy of OAT LIT DA		e, or non naza		Designated Drop Ship			inature:					
	number of page 2.					500 p. 0 101		<i></i>	01	,					

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

Version 2024 For Designa	ted Drop Ship Only Products, Please Use Page 3				
MATERIAL HA	ZARD CLASSIFICATION and TRANSPORTATION				
Is this product (check all that apply): a. Cytotoxic? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? No Is the product a CA Prop 65 reproductive toxicant? 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? Is this product regulated for shipment by DOT? No	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				
(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 the product a NIOSH hazardous drug? If yes, indicate which: Hazardous Waste Identification EPA Hazardous Waste Code: Waste Characteristics				
Is this product regulated for shipment by IATA? No (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? No If Yes, is it managed with a pharmacy registry? No Website URL: Image: Colored and Col				
Is the product restricted for air shipment? If so, indicate restriction: Passenger No Cargo Passenger & Cargo Is this a reactable quantity? No	Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?) REMS: No				
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 Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	REMS: No REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Phone: Wholesale distributor support: Provider Name: Site Enrollment Number assigned DEA #: by Supplier: NPI #:				
SP#	Registry: No Registry Program Contact Name: Phone:				
ADD'L STORAGE INFORMATION Is the Product Controlled Substance Code	Comments RETURN INSTRUCTIONS				
Controlled Substance? No Controlled Substance Code Controlled by State(s)? No Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which: No Schedule No. Is it a scheduled listed chemical product?: No	KEI URN INSTRUCTIONS Contact tel. # if product received damaged: 1-866-827-3647 Is product returnable for credit: Yes URL/Link to returns policy: Yes				
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	contact - customerservice@camberpharma.com				
Restricted to retail pharmacy only: No 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? No If so, which states? Other requirements? Comments?				
MISCELLANE	OUS NOTES and/or Image of Product Barcode:				



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Version 2024	FOR DESIGNATED DROP SHIP PRODUCT ONLY - if r	ot a designated drop ship, do not complete.	
Order Method for Des	signated 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 Expedited Freight Charges o		Purchase order daily receipt cut off time by supplier Cut off time: Shipping lead time of PO: Hours Da Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt: Overnight and Priority Overnight PO Processing	ays
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:		Days of week overnight is available: Monday	
Comments:	s only:	Days of week overnight is available: Tuesday Tuesday Wednessi Priority Overnight receipt available: Friday PO Receipt Cut off time: PO Receipt Cut off time: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Policity	/ day
Other Data Informati	ion 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?	
Miscell	aneous Notes:		
		ADDITIONAL INFORMATION	
		Is product order for scheduled patient procedure? Is product order for restocking purposes?	