

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

Version 2024						Introduction Typ	De: New Item	х	Final Version			Date:	7/15	/2024
			PRODUCT INFORMAT	ION					SPECIAL HAN	NDLING AND STOR	AGE REQUI	REMENTS*		
Company Name:	Camber Pharma	ceuticals, Inc.				Applicatio	n: ANDA	a. Temperature – Inc	dicate the USP temp	erature range for th	nis product.			
Application Number for NDA/AND		0(k): 205504	4			NDA 505(b) Type:	NOT APPLICABLE	Temp	erature Range	Controlled Room -	- between 20	and 25 C (68	° – 77° F)	
Medical Device Class, if applicable:														
DUNS:	11-856-3719								Temperature Range	Requirement				
Proprietary Name (If Applicable) ar Selling Unit NDC:	31722-857-30	ame: Eszopi	clone Tablets, USP 3 mg Unit of Use NDC:		31722-857-30	UPC: 3	31722857307	Notes	(write in)					
UDI	31722-037-30		CVX Code:		31722-037-30	MVX Code:	31722037307	Notes						
	Eszopiclone Tabl	loto LICD 2 mg						lo thio	product to be shippe	d to quotomoro on ic	202		No	1
Description:	ESZOPICIONE TADI	iets, OSF 3 mg							product to be shippe				No	-
Active Ingredient(s):		Eszopiclone, USP						1	p		.,			1
								b. Contact for temper		estions:				
URL for Additional Product Informa		www.camberpharma	a.com					Name			Soma Raju	_		
	800 Centennial A Piscataway	Ave, Suite 1			State:	Address 2:	Zip: 08854	Numb			732-529-042			
	Customer Service	Α			Email:	customerservice@c		Group E-mail: somaraju@heterousa.com						
Phone Number:	1-866-827-3647				Fax:	732-562-8788		c. Special regulation	s for product in any	states?			*Yes	1
Product Therapeutic Classification	1:	Sedative-hypnotic			1				al returns requiremen				No	1
_								_						_
	ADDIT	IONAL PRODUCT IN	FORMATION			PRODUCT DE	SCRIPTION INFORMATION	d. Store product (un	it of sale) upright?				No	
The product is?			Is the Product	Direct-Ship C	only				ct product (unit of s	ale) from light?			No	
a legend device?		No	Is the Product	Unit of Use		Size:	30 ct	e. Shelf life:					24	Months
if yes, enter class # a product kit?		No	Orphan Drug Status				3 mg	Initial	shelf life at launch	(if different):				Months
if yes, list NDCs of		INO	FDA Approval Status			Strength:	3 mg			ORDER INFORM	IATION			
component parts			. D			Dosage Form:	Film-coated tablet							
reverse numbered?		No				Dosage Form:			of Sale		What is the		unit?	
co-licensed?		No	Allergens Present					x			1 Bottle of 3			
latex-free? preservative-free?		Yes	Dairy, Lact	ose, Casein		Product Shape	Round, biconvex		Box/Carton		(Write-in, e.	g. 1 Box of 1	J Vials)	
correctional institution block?		Yes No					Dark blue to blue		Ampule Glass		Minimum or	der quantity	17	Yes
opioid?		No				Product Color:	Bain side to side		Tube			uoi quaiitity	•	
Cannabinoid?		No	Country of Origin	India		Product Imprir	Debossed with 'H' on one side and 'E16' on the other side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to un	nit dose for					T Toddot IIIIpili	E 16 Off the other side		Vial Liquid Multi				ich package	type?
hospital scanning? If Unit Dose, indicate NDC here:			Is this product covered u Trade Agreements Act (T		No				Vial Powder Sgl Vial Powder Multi		24	Each Inner/Carton	·/Deels	
Il Onit Dose, indicate NDC here:			Trade Agreements Act (1	AA)!	INO				Other: Write In			Case	/Pack	
			FOR GENERIC DRUG PRO	DDUCTS										
											-			
					Au		If Authorized Generic, other		Pl	HARMACY ORDER	/ BILL UNIT			
	AB					s	ection fields are not applicable	Rec. sell unit to cus	tomer?	-	Rx billing u		acy:	
II. Generic Equivalent to What Brand?: Lunesta							Each Gram							
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION (Write-in, e.g. 1 Vial) Gram HCPCS J-Code: Milliliter														
			•											
Does supplier meet DSCSA definit	tion of manufactu	irer?	Yes		GLN:	860000397957			ITE	M AND PACKING IN	NFORMATION	١		
Is product exempt from DSCSA?			No											
If yes, select exemption:					GCP:				Weight Lbs.		ons (US msm	•	Volume	Saleable #
Other exemption - Write in: Is product repackaged?			No		If yes was or	riginal product purch	asad	Item/Each:		Depth	Width	Height	(Cube)	Pieces
Is product sold by manufacturer's	exclusive distrib	utor?	Yes		direct from m		uoou	inong Euroni	0.05	1.5	1.5	3	6.75	1
Has FDA granted waiver/exception	n/exemption for p		No		Provide sour	ce manufacturer for I	repackaged product	Box/Carton/Bundle/						
If yes, attach documentation from	n FDA.							Inner Pack:						
		GTII	N AND HIBCC PRODUCT IN	IFORMATION.				Case:	1.8	9.5	6.5	4	247	24
		011	I AND HIBOUT RODUCT II	ORMATION				Pallet:						
Saleable Unit of Measure	RFID tag(Y/N)	Saleable	HIBCC		GTI	N-14	Unit of Use GTIN-14							
		Quantity			_									
x Item/Each	N	1			003	31722857307	00331722857307		OST INFORMATION			MUOLECAL	ER USE ONL	V.
Box/Carton/Bundle/Inner Pack X Case	N	24			303	31722857308		C	JST INFORMATION			WHOLESALI	ER USE UNL	.T:
Pallet	.,	24			303	322007000		Regular Cost			Vendor #:			
								Invoice Cost (WAC)	(\$)	\$5.25	Whsl. Code			
											Fineline Co	de:		
								As of date:	6/12/2024		Į.			
			Attach copy of SAFETY DA	TA SHEET (SF	S) or non haza	rd letter, PACKAGE IN	NSERT, LABEL AND PHOTO OF	PRODUCT PACKAGING	and BARCODE.		1			
*Please provide any additional info	ormation on page	2.			.,		esignated Drop Ship Only.	Signa						



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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? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant?	SDS Hazard Classification						
Is the product a CA Prop 65 carcinogen? Is the product a CA Prop 65 reproductive toxicant? No Does the product label bear a CA Prop 65 warning? No	x Organic Corrosive Inorganic 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? No	Does the product have an Aerosol class? If yes, 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?	El A Hazardous Waste Oode.						
(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: NO Phone: DEA #: NCPDP#: NCPDP#: NPI #:						
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	Comments						
SP#	Registry: Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
Is the Product Controlled Substance? Controlled Substance Code Controlled by State(s)? Yes Controlled Substance Code Listed Chemical (List I or II) No	RETURN INSTRUCTIONS						
ARCOS Reportable? Schedule No. Yes If yes, indicate which: Is it a scheduled listed chemical product?: No CLASS OF TRADE RESTRICTION:	Contact tel. # if product received damaged: Is product returnable for credit: URL/Link to returns policy: 1-866-827-3647 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	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:							
	EOUS NOTES and/or Image of Product Barcode:						
*Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR Pa	rt 1301.72.						



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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 Phone No.:	Shipping lead time of PO: Hours Days						
e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Name: Phone:	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:	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/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?						