

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

Version 2024						Introduction Ty	vpe: New Item		x Final Version			Date:	6/10	/2024
			PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOR	AGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc.						Applicati	on: ANDA	a. Temperature – Indicate the USP temperature range for this product.						
Application Number for NDA/AN	IDA/BLA; PMA/510	0(k):	211228			NDA 505(b) Type:	NOT APPLICABLE		emperature Range	Controlled Room		and 25 C (68	3° – 77° F)	
Medical Device Class, if applica	ble:													
DUNS:	11-856-3719							Ot	her Temperature Range F	Requirement				
Proprietary Name (If Applicable) a		ame:	Methadone Hydrochloride Tablets,						(write in)					
Selling Unit NDC:	31722-947-01		Unit of Use NDC:				331722947015	No	otes					
UDI			CVX Code:			MVX Code:								
Description:	Methadone Hydro	ochloride Tablet	ts, USP 10 mg						this product to be shipped				No	
								Is	this product to be shipped	d to customers on o	Iry ice?		No	
Active Ingredient(s): Methadone hydrochloride, USP														
URL for Additional Product Information: www.camberpharma.com								b. Contact for temperature excursion questions: Name: Soma Raju						
Address:	800 Centennial A		priama.com		T	Address 2:			umber:		732-529-042	23		
City:	Piscataway	,			State:	NJ	Zip: 08854	Group E-mail: somaraju@heterousa.com						
Key Contact:	Customer Service	е			Email:	customerservice@	camberpharma.com							
Phone Number:	1-866-827-3647				Fax: 732-562-8788			c. Special regula	*Yes					
Product Therapeutic Classification	on:	Opioid agonis	st					Special returns requirements for this product?					*Yes	
					_									_
	ADDITI	IONAL PRODU	ICT INFORMATION			PRODUCT D	ESCRIPTION INFORMATION	d. Store product	(unit of sale) upright?				No	
The product is?			Is the Product	Direct-Ship (Only				otect product (unit of sa	ale) from light?			No	
a legend device?		No	Is the Product	Neither		Size:	100 ct	e. Shelf life:					24	Months
if yes, enter class #		Ta a	Orphan Drug Status					In	itial shelf life at launch (if different):				Months
a product kit?		No	FD 4 4 01-11-1			Strength:	10 mg			ORDER INFORM	MATION			
if yes, list NDCs of component parts			FDA Approval Status				Tablet			OKDEK INFORM	IATION			
reverse numbered?		No				Dosage Form	: Tablet	ll u	nit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present				L	'II	x Bottle		1 Bottle of 1			
latex-free?		Yes		Alcohol		Product Shap	Round, beveled edge		Box/Carton		(Write-in, e.	.g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Corn,	Alconoi		Product Snap			Ampule					
correctional institution block?		No				Product Color	White to off-white		Glass		Minimum o	rder quantity	1?	Yes
opioid?		Yes							Tube					
Cannabinoid?		No	Country of Origin	USA		Product Impri	nt: Scored on one side and debossed 'T293' on the other side		Vial Liquid Sgl		W. V 1			
If Unit Dose, is item bar coded to unhospital scanning?	unit dose for		Is this product covered u	under the					Vial Liquid Multi Vial Powder Sql			many of whi	icn package	type?
If Unit Dose, indicate NDC here:			Trade Agreements Act (Yes				Vial Powder Multi		24	Inner/Cartor	/Pack	
iii ciiii 2000, iiialoalo 1120 11010.				,.	. 00				Other: Write In			Case	W GOIL	
			FOR GENERIC DRUG PR	ODUCTS								-		
					Au	uthorized Generic	*If Authorized Generic, other		PH	IARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AA						section fields are not applicable	Rec. sell unit to customer? Rx billing unit to pharmacy:						
II. Generic Equivalent to What Brand?: Dolophine						Each								
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFOR				PMATION			(Write-in, e.g. 1 Vial) HCPCS J-Code: Gram Milliliter							
		DRUG	SOFFET CHAIN SECURITY ACT	DSCSA) INFO	KIWIATION			HCPCS J-Code:		1		Milliliter		
Does supplier meet DSCSA defini	ition of manufactu	irer?	Yes	_	GLN:	0860000397957			ITEN	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			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		If yes, was or	riginal product purcl	hased	Item/Each:	0.07	1.52	1.52	2.69	6	1
Is product sold by manufacturer's			Yes		direct from m					1.02	1.52	2.03	0	· '
Has FDA granted waiver/exceptio		roduct?	No		Provide sour	ce manufacturer for	repackaged product	Box/Carton/Bund	dle/					
If yes, attach documentation fro	m FDA.							Inner Pack: Case:						
			GTIN AND HIBCC PRODUCT I	NFORMATION				Case:	2.1	10.8	7	3.2	242	24
								Pallet:						
Saleable Unit of Measure	RFID tag(Y/N)	Saleable	HIBCC		GTI	IN-14	Unit of Use GTIN-14							
		Quantity												
x Item/Each	N	1			003	331722947015			OCCUPATION A TION			MILOL FO	ER USE ONL	V-
Box/Carton/Bundle/Inner Pack		- 04			100	04700047040			COST INFORMATION			WHOLESAL	ER USE ONL	_Y:
X Case Pallet	N	24			103	331722947012		Regular Cost			Vendor #:			
Pallet		-						Invoice Cost (WA	AC) (\$)	\$16.00	Whsl. Code	#-		
									/ \+/	ψ10.00	Fineline Co			
								As of date:	4/29/2019					
11											1			
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For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HA	IZARD CLASSIFICATION and TRANSPORTATION						
Is this product (check all that apply):							
a. Cytotoxic?	SDS Hazard Classification						
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?							
Is the product a CA Prop 65 carcinogen?	x Organic Corrosive						
Is the product a CA Prop 65 reproductive toxicant?	Inorganic Oxidizer						
Does the product label bear a CA Prop 65 warning?	Steroid/Androgen Contact Hazard						
,							
c. Contact Hazard?	Does the product have an Aerosol class? If yes, No						
d. Does this product require special clean-up instructions?	identify NFPA Storage Level:						
(If yes, attach SDS with special instructions.)	NFPA Storage Level:						
e. Does the product contain DEHP?							
	In the conduct of NICOU become drawn						
Is this product regulated for shipment by DOT? No	Is the product a NIOSH hazardous drug? No If yes, indicate which:						
(if yes, answer a-e below and provide SDS) a. UN/Identification Number	ii yes, indicate wilicit.						
b. Proper Shipping Name							
c. DOT Hazard Class	Hazardous Waste Identification						
d. Packing Group	TRADE TO THE STATE OF THE STATE						
e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics						
	1.11.11.11.11.11.11.11.11.11.11.11.11.1						
Is this product regulated for shipment by IATA?	REMA REGISTRY DESTRUCTIONS						
(if yes, answer a-e below and provide SDS)	REMS or REGISTRY RESTRICTIONS						
a. UN/Identification Number							
b. Proper Shipping Name	Is there a REMS on this product?						
c. DOT Hazard Class	If Yes, is it managed with a pharmacy registry?						
d. Packing Group	Website URL: https://opioidanalgesicrems.com/home.html						
e. Inhalation Hazard?							
Is the product restricted for air shipment? If so, indicate restriction:	Med Guide Required Yes						
Passenger	Limited Distribution Requirement No						
Cargo	Comments / Details: (For example, iPledge program?)						
Passenger & Cargo							
Is this a reportable quantity? No	REMS: Yes						
RQ Threshold:	REMS Program Manager Name: Prathima Arrabelly Phone: (631) 881-4614 Ext. 1412						
Is this a marine pollutant? No	Supplier Manages REMS registry exclusively:						
Is this product shipped utilizing an authorized DOT exception or Special Permit?	Wholesale distributor support:						
No (if yes, identify method below)	Provider Name: DEA #:						
Limited Quantity	Site Enrollment Number assigned NCPDP#:						
Consumer Commodity, ORM-D	by Supplier: NPI #:						
Small Quantity (49 CFR 173.4)							
Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101);							
SP#	Registry: No No						
	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
Is the Product							
Controlled Substance? Yes Controlled Substance Code 9250	RETURN INSTRUCTIONS						
Controlled by State(s)? Yes Listed Chemical (List I or II) No							
ARCOS Reportable? Yes If yes, indicate which:	Contact tel. # if product received damaged: 1-866-827-3647						
Schedule No. 2 Is it a scheduled listed chemical product?: No	Is product returnable for credit:						
CLASS OF TRADE RESTRICTION:	URL/Link to returns policy:						
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	contact - customerservice@camberpharma.com						
Restricted to retail pharmacy only:	Special regulations or returns requirements for this						
Restricted to hospital, clinics, and physician offices only:	product in certain states? Yes						
Restricted from US territories? (explain in comments)	If so, which states? Other requirements? Comments?						
Comments:	DEA Form 222 or its electronic equivalent is required for all returns in all states.						
MOOFILANG	CIUS NOTES and lay Image of Product Parcedo						
	OUS NOTES and/or Image of Product Barcode:						
*Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR Part	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 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: PO Receipt cut off time: Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday					
	Priority Overnight receipt available:					
Class of Trade Restriction: 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:	PO Receipt Cut off time: Saturday Overnight receipt available: PO Receipt Cut off time: Phone: 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: Miscellaneous Notes:	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?					
	ADDITIONAL INFORMATION					
	Is product order for scheduled patient procedure? Is product order for restocking purposes?					