

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

Version 2021					Introduction 1	Туре:	New Item]	x Final Version			Date:	6/4/:	2024		
PRODUCT INFORMATION								AGE REQUI	REMENTS*							
Company Name: Camber Pharmaceuticals, Inc.				Application: ANDA			a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 215523 a. leniperature remains continuation of the properature Range and the properature Range (Controlled Room – between 20 and 25 C (68° – 77° F)																
Medical Device Class, if applicab				7.							romporataro rtango	-				
DUNS:	11-856-3719									1	Other Temperature Range	Requirement	Evoursions	permitted betw	veen 15°C a	nd 30°C
Proprietary Name (If Applicable) a		ame.	Devmeth	nylphenidate Hydrochloride	Evtended-Rele	ase Cansules	20 ma			1	(write in)	requirement	(59°F to 86°		Jeen 15 C al	11d 30 C
	31722-232-01	ume.	DOMINION	Unit of Use NDC:	Exteriaca recie	asc Capsuics	UPC:	331722	232012	1	Notes		(00 : 10 00	,		
UDI				CVX Code:			MVX Code:	001122		1	110100					
										1						
Description:	Dexmethylphenida	ate Hydrochio	oride Exten	nded-Release Capsules 20	mg						Is this product to be shipped				No	
A - 11 I 111(-)		Danier attacks	de a setala da de	o de esta esta esta esta esta esta esta est							Is this product to be shipped	to customers on	ary ice?	L	No	
Active Ingredient(s):		Dexmetnyip	onenidate r	hydrochloride												
URL for Additional Product Inform	nation:	waww cambe	ornharma	com						b. Contact for	r temperature excursion qu Name:	estions:	Soma Raju			
Address:	htion: www.camberpharma.com 300 Centennial Ave, Suite 1				Address 2:			-	Number:			732-529-0423				
City:	Piscataway				State:		NJ Zip : 08854			Group E-mail:			somaraju@heterousa.com			
Key Contact:	Customer Service					customerservice			Situation of the state of the s							
Phone Number:	1-866-827-3647				Fax:	732-562-8788	Courne	ipnama.com	c. Special regulations for product in any states?					*Yes		
Product Therapeutic Classification					· un	102 002 0100										
1 Toddet Therapeutic Glassification	Classification: Central nervous system (CNS) stimulant Special returns requirements for this product? *Yes															
	ADDITIO	ONAL PRODU	UCT INFO	RMATION			PRODUCT	DESCRIE	TION INFORMATION	d Store prod	uct (unit of sale) upright?			Г	No	
The weeduction	7,551110			Is the Product	Direct-Ship O	inly	- 1.05001			1		ala) fram !!!- c		L	No	
The product is?		NI-	7		Neither	riiy		T.	100 ct	e. Shelf life:	Protect product (unit of s	ale) from light?			24	Mantha
a legend device?		No		Is the Product	Neitriei		Size:		100 ct	e. Sheir lire:	Initial abolf life at laurah f	if different).			24	Months
if yes, enter class #		INI-		Orphan Drug Status					20		Initial shelf life at launch (ir airierent):		Į.		Months
a product kit? if yes, list NDCs of		No		FDA Approval Status	A A				20 mg		ORDER INFORMATION					
component parts				FDA Approvai Status					Extended-release, hard			ORDER IN OR	IATION			
reverse numbered?		No					Dosage Form		gelatin capsule		Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	-	Allergens Present				3	gelatii oapsale		x Bottle		1 Bottle of 1		unit.	
latex-free?		Yes	-					(Capsule		Box/Carton			g. 1 Box of 10) Vials)	
preservative-free?		Yes	-	Corn, Alcohol, An	imal, Sugar, D	ye	Product Sha	ape:	oapsuic		Ampule		(vviite iii, e	g. I Dox of To	viais)	
correctional institution block?		No							Light brown cap and		Glass		Minimum o	rder quantity	12	Yes
opioid?		No					Product Col		white opaque body		Tube			ao. quantity		.00
Cannabinoid?		No	-	Country of Origin	USA			ī	mprinted with 'M20' on cap		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	nit dose for	1.10	_	,			Product Imp	orint:	and 'AC' on body in black ink		Vial Liquid Multi		If Yes, how	many of which	ch package	type?
hospital scanning?				Is this product covered un-	der the						Vial Powder Sgl			Each	р	.,,,,,
If Unit Dose, indicate NDC here:				Trade Agreements Act (TA		Yes					Vial Powder Multi			Inner/Carton/	/Pack	
·											Other: Write In			Case		
FOR GENERIC DRUG PRODUCTS																
						Au	thorized Generic		orized Generic, other		PH	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB						section fields are not applicable			Rec. sell unit	Rx billing unit to pharmacy:					
II. Generic Equivalent to What Bra	ind?:	Focalin XR	Focalin XR								Each					
								(Write-in, e.g. 1 Vial)				•	Gram			
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION																
Does supplier meet DSCSA definit	tion of manufactu	rer?		Yes No	-	GLN:	0860000397957				ITEM	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?				NO												
If yes, select exemption:						GCP:					Weight Lbs.	Dimensi	ons (US msn	•	Volume	Saleable #
Other exemption - Write in:												Depth	Width	Height	(Cube)	Pieces
Is product repackaged?				No			riginal product			Item/Each:	0.15	2.1	2.1	3.50	15.44	1
Is product sold by manufacturer's				Yes			irect from mfr?								-	
Has FDA granted waiver/exception		roduct?		No		Provide sour	ce manufacturer f	or repac	kaged product	Box/Carton/B	undle/					
If yes, attach documentation from	m FDA.									Inner Pack:						
			GTIN A	AND HIBCC PRODUCT INF	OPMATION					Case:	4.2	13.25	9	4.5	536.63	24
			GIIIVA	AND THE CCT RODOCT IN	OKMATION					Pallet:						
Saleable Unit of Measure	S	Saleable Quan	ntity	HIBCC		GTI	N-14		Unit of Use GTIN-14	I dilet.						
X Item/Each	_	1	1				31722232012	T								
Box/Carton/Bundle/Inner Pack										COST INFORMATION			WHOLESALER USE ONLY:			
x Case		24				103	31722232019									
Pallet										Regular Cost			Vendor #:			
	Ī							T		Invoice Cost		\$132.00	Whsl. Code	#:		
	I									11			Fineline Co			
	I									As of date:	1/13/2022					
	I									[]						
										Ш						
		_	At	tach copy of SAFETY DATA	A SHEET (SDS	6) or non hazaı				PRODUCT PACK	AGING and BARCODE.					
*Please provide any additional info	ormation on page	2.					See new p. 3 for	r Desian:	ated Drop Ship Only.		Signature:					



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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?	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?							
Is this product regulated for shipment by DOT?	Is the product a NIOSH hazardous drug?						
(if yes, answer a-e below and provide SDS)	If yes, indicate which:						
a. UN/Identification Number							
b. Proper Shipping Name							
c. DOT Hazard Class	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)	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:						
e. Inhalation Hazard?							
Is the product restricted for air shipment? If so, indicate restriction:	Med Guide Required No						
Passenger	Limited Distribution Requirement						
Cargo	Comments / Details: (For example, iPledge program?)						
Passenger & Cargo							
Is this a reportable quantity? No	REMS: No						
RQ Threshold:	REMS Program Manager Name: Phone:						
Is this a marine pollutant? No	Supplier Manages REMS registry exclusively:						
Is this product shipped utilizing an authorized DOT exception or Special Permit? No (if yes, identify method below)	Wholesale distributor support: Provider Name: DEA #:						
No (if yes, identify method below) Limited Quantity	Site Enrollment Number assigned NCPDP#:						
Consumer Commodity, ORM-D	by Supplier: NPI #:						
Small Quantity (49 CFR 173.4)	бу барыст.						
Special Permit; DOT-SP	Comments						
Special Provision (listed in Column 7 of 49 CFR 172.101);							
SP#	Registry: No						
	Registry Program Contact Name: Phone:						
ADD'L STORAGE INFORMATION	Comments						
Is the Product							
Controlled Substance? Yes Controlled Substance Code 1724	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:						
	contact - customerservice@camberpharma.com						
	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:	product in certain states?						
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.						
MISCELLAN	EOUS NOTES and/or Image of Product Barcode:						
*Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR F	art 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 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?