

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

Version 2021						Introduction Type:	New Item		x Final Version			Date:	6/4/2	/2024
			PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOR	RAGE 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): 214959							Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)							
Medical Device Class, if applicable:														
DUNS:	11-856-3719							l l	Other Temperature Range F	Requirement	Excursions i	permitted to 1	5° to 30°C (5	59° to 86°F)
Proprietary Name (If Applicable) a		me: Dextroampl	hetamine Saccharate, Amphetamine Aspartate Monoh	drate, Dextroamphetamine Su	fate, and Amphetamine Sulfa	ate Extended-Release Capsules (Mixed Salts of	a Single Entity Amphetamine Product) 20 mg	ī	(write in)	toquiromont	ZXOUIOIOIIO	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	0,000,0	, ,
Selling Unit NDC:	31722-188-01		Unit of Use NDC:			UPC: 331	722188012	1	Notes					
UDI			CVX Code:			MVX Code:								
Decementary	Dautraamahatamia	a Casabarata Ama		dente Deutrone	- h - t : C   f		ulfate Cutended Delega	i	la thia anadust ta ha ahiana.		2		No	
Description:			hetamine Aspartate Monohy		prietamine Suii	ate, and Amphetamine 5	uliale Extended-Release		Is this product to be shipped Is this product to be shipped					-
Capsules (Mixed Salts of a Single Entity Amphetamine Product) 20 mg								·	is this product to be shipped	i to customers on t	aly ice :		No	_
Active Ingredient(s):  Dextroamphetamine saccharate, amphetamine aspartate monohydrate, dextroamphetamine sulfate, USP, and amphetamine sulfate, USP								h Contact for	r temperature excursion que	etione:				
URL for Additional Product Information: www.camberpharma.com								b. Contact io	Name:	sations.	Soma Raju			
Address:	800 Centennial Av					Address 2:			Number:		732-529-042	23		
City:	Piscataway					o: 08854		Group E-mail:			neterousa.cor	m		
Key Contact:	Customer Service				Email:	customerservice@can							_	
Phone Number:	1-866-827-3647				Fax:	732-562-8788		c. Special reg	gulations for product in any	states?			*Yes	1
Product Therapeutic Classification	n:	Central nervous sy	stem (CNS) stimulant		1			'  '	Special returns requirement				*Yes	1
			, ,		1				-,					_
	ADDITIO	NAL PRODUCT IN	IFORMATION			PRODUCT DESC	CRIPTION INFORMATION	d. Store prod	uct (unit of sale) upright?				No	7
The product is?			Is the Product	Direct-Ship C	nly			11	Protect product (unit of sa	le) from light?			No	ī
a legend device?		No	Is the Product	Neither	,	1	100 ct	e. Shelf life:	Stoot product (unit 0) So	,			24	Months
if yes, enter class #			Orphan Drug Status			Size:	.00 00	S. Onen me.	Initial shelf life at launch (	f different).			4-7	Months
a product kit?		No	Orphan Drug Otatus				20 mg		initial shell life at launen (	r diricicity.				_ months
if yes, list NDCs of		110	FDA Approval Status			Strength:	g			ORDER INFOR	MATION			
component parts			, ,				Extended-release, hard							
reverse numbered?		No				Dosage Form:	gelatin capsule		Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present				-		x Bottle		1 Bottle of 1	00 Capsules		
latex-free?		Yes	Dye, Corn, Alcol	ad Animal Cu		Product Shape:	Capsule		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Dye, Com, Alcon	ioi, Aililiai, Su	yaı	Froduct Snape.			Ampule					
correctional institution block?		No				Product Color:	Orange opaque cap and		Glass		Minimum o	der quantity	/?	Yes
opioid?		No				rioduct color.	orange transparent body		Tube					
Cannabinoid?		No	Country of Origin	USA		Product Imprint:	Imprinted with '20 mg' on cap		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for						and 'T' on body in black ink		Vial Liquid Multi				ich package i	type?
hospital scanning?			Is this product covered u						Vial Powder Sgl		24	Each		
If Unit Dose, indicate NDC here:			Trade Agreements Act (	ΓAA)?	Yes				Vial Powder Multi			Inner/Cartor	n/Pack	
									Other: Write In			Case		
			FOR GENERIC DRUG PR	ODUCTS										
					A	ubasinad Casasia *If A			BU	ARMACY ORDER	/ DILL LINIT			
				_	Au		Authorized Generic, other tion fields are not applicable	_		ARMACT ORDER				
I. Orange Book Rating:	AB1					360	tion helds are not applicable	Rec. sell unit	to customer?	1	Rx billing u		acy:	
II. Generic Equivalent to What Bra	ind?:	Adderall XR						0.000	4 \ / / - D			Each		
		DRIIG SIIDDI	LY CHAIN SECURITY ACT (	DSCSA) INFOE	PMATION			(Write-in, e.g.	. 1 viai)			Gram Milliliter		
		DRUG SUFFI	ET CHAIN SECONTT ACT	DSCSA) IN OF	MATION							wiiiiiitei		
Does supplier meet DSCSA definit	tion of manufacture	er?	Yes		GLN:	0860000397957			ITEN	AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?		· ·	No											
If ves. select exemption:					GCP:			i I		Dimens	ions (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:					JOI .			' <b> </b>	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No		If ves, was or	riginal product purchase	ed	Item/Each:						
Is product sold by manufacturer's	exclusive distribut	tor?	Yes		direct from m				0.13	2.01	2.01	3.4	13.74	1
Has FDA granted waiver/exception			No	7	Provide sour	ce manufacturer for rep	ackaged product	Box/Carton/B	Bundle/					
If yes, attach documentation fror	m FDA.							Inner Pack:						
								Case:	3.4	12.3	8.3	3.8	387.94	24
		GT	IN AND HIBCC PRODUCT I	NFORMATION					0.1	12.0	0.0	0.0	007.07	
								Pallet:						
Saleable Unit of Measure	Sa	aleable Quantity	HIBCC			N-14	Unit of Use GTIN-14							
X Item/Each		1			003	31722188012			COST INCODMA <del>TION</del>		_	MUOLECAL	ER USE ONL	I V.
Box/Carton/Bundle/Inner Pack		24			100	24722400042			COST INFORMATION			WHOLESAL	ER USE ONL	3.5
X Case		24			103	31722188019		Bamulas Care			Vendor #:			
Pallet					-			Regular Cost		070.00		4.		
	-							Invoice Cost	(AAWC) (9)	\$70.00	Whsl. Code			
	-							As of date:	10/25/2021		Fineline Co	uc.		
	-							As or date:	10/20/2021		-			
	_							l I						
			Attach copy of SAFETY DA	ATA SHEET (SD	S) or non haza	ard letter, PACKAGE INSI	ERT, LABEL AND PHOTO OF F	RODUCT PACK	AGING and BARCODE.		1			



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

#### Version 2021

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?  No  Does the product label bear a CA Prop 65 warning?  No	Inorganic Oxidizer Steroid/Androgen Contact Hazard						
boes the product laber bear a CA Frop 65 warning!	Sterotovariologen						
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? No						
(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	Nuzurodas Waste Identification						
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 d. Packing Group	If Yes, is it managed with a pharmacy registry?  Website URL:						
e. Inhalation Hazard?	Website Orc.						
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 #:						
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						
ADD'L STORAGE INFORMATION	Registry Program Contact Name: Phone:						
	Comments						
Is the Product  Controlled Substance?  Yes  Controlled Substance Code 1100	RETURN INSTRUCTIONS						
Controlled by State(s)? Yes Listed Chemical (List I or II) No	KETOKKINGTIGGTIGKE						
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?						
Restricted from US territories? (explain in comments)  No	If so, which states? Other requirements? Comments?						
Comments:	DEA Form 222 or its electronic equivalent is required for all returns in all states.						
	·						
MISCELLANE	OUS NOTES and/or Image of Product Barcode:						
*Storage of this product must abide by the federally mandated DEA requirements outlined in 21 CFR Par							
Storage of this product must able by the redefany manualed DEA requirements outlined III 21 OFN Fall	100 1.1 2.						



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#### FOR DESIGNATED DROP SHIP PRODUCT ONLY - if not a designated drop ship, do not complete.

Order Method for Designated Drop S	nip Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI		Purchase order daily receipt cut off time by supplier Cut off time:
b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	per:	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 Designa	ed Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order:  Drop Ship service fee billed with each order:		Overnight receipt available:  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, clinic Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	s and physician offices	Saturday Overnight receipt available:  PO Receipt Cut off time:  Phone: Fax: EDI:  Overnight Fees apply: Other fees apply:
Other Data Information Required to F	rocess 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?