

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

Version 2021						Introduction	Туре:	New Item		x Final Version			Date:	10/5/	/2023	
			PRODUCT INFORMAT	ION						SPECIAL HA	NDLING AND STO	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): 214956 214956 214956 214956																
Medical Device Class, if applical	ole:								1							
DUNS:	11-856-3719									Other Temperature Rang	e Requirement		permitted bet	ween 15°C to	30°C (59°F	
Proprietary Name (If Applicable) a		me: Gab	papentin Capsules, USP 300 mg	9						(write in)		to 86°F)				
Selling Unit NDC:	31722-149-05		Unit of Use NDC:			UPC:	33172	2149051	-	Notes						
UDI			CVX Code:			MVX Code:			-							
Description:	Gabapentin Capsul	les, USP 300 mg								Is this product to be shipp				No		
Is this product to be shipped to customers on dry ice? No																
Active Ingredient(s): Gabapentin b. Contact for temperature excursion questions:																
URL for Additional Product Information: www.camberpharma.com								b. Contact 10	Name:	questions.	Soma Raju					
Address:	1031 Centennial Ave (and) 800 Centennial Ave, Suite 1					Address 2:							732-529-0423			
City:				State:	NJ Zip: 08854			Group E-mail:			somaraju@heterousa.com					
Key Contact:	Customer Service	ustomer Service			Email:		ce@camberpharma.com									
Phone Number:	1-866-827-3647				Fax:	732-562-8788			c. Special regulations for product in any states?			*Yes				
Product Therapeutic Classificatio	n:	Anticonvulsant								Special returns requireme	ents for this product	?		No		
	ADDITIO	NAL PROPUSE	NIEGOMATION .			2222127	D=000	DTION INFORMATION								
	ADDITIO	NAL PRODUCT				PRODUCT	DESCRI	PTION INFORMATION	d. Store prod	uct (unit of sale) upright?				No		
The product is?			Is the Product	Direct-Ship C	Only					Protect product (unit of	sale) from light?			No		
a legend device?		No	Is the Product	Neither		Size:		500 ct	e. Shelf life:					24	Months	
if yes, enter class # a product kit?		No	Orphan Drug Status					000		Initial shelf life at launch	if different):				Months	
if yes, list NDCs of		INO	FDA Approval Status			Strength:	300 mg	ORDER INFORMATION				TON				
component parts			1 DA Approvar otatus					Hard gelatin capsule								
reverse numbered?		No				Dosage For	m:			Unit of Sale		What is the	NDC selling	g unit?		
co-licensed?		No	Allergens Present							x Bottle		1 Bottle of 5	00 Capsules			
latex-free?		Yes	Corn, Alcohol, Ar	imal. Wheat. [Ove	Product Sha	ape:	Capsule		Box/Carton		(Write-in, e	.g. 1 Box of 1	0 Vials)		
preservative-free?		Yes		,,	,-					Ampule						
correctional institution block?		No				Product Col	lor:	Yellow opaque cap and		Glass		Minimum o	rder quantit	y?	Yes	
opioid? Cannabinoid?		No No	Country of Origin	USA				yellow opaque body 'A' on cap and '470' on		Tube Vial Liquid Sgl						
If Unit Dose, is item bar coded to u		INO	Country of Origin	USA		Product Imp	print:	body		Vial Liquid Sgi		If Yes how	many of wh	ich package	tyne?	
hospital scanning?	init dosc for		Is this product covered ur	nder the				,		Vial Powder Sq			Each	ion package	турс.	
If Unit Dose, indicate NDC here:			Trade Agreements Act (T		Yes					Vial Power Mult			Inner/Cartor	n/Pack		
										Other: Write In			Case			
			FOR GENERIC DRUG PRO	DUCTS												
Authorize					thorized Generic				F	HARMACY ORDER	R / BILL UNIT					
I. Orange Book Rating:	AB					section fields are not applicable			Rec. sell unit to customer?			Rx billing unit to pharmacy:				
II. Generic Equivalent to What Bra	and?:	Neurontin Capsu	lle							Each						
			N V 01141N 05011DIEV 407 ((Write-in, e.g. 1 Vial)							
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION								Milliliter								
Does supplier meet DSCSA defini	tion of manufactur	or?	Yes	7	GLN:	031722000000				ITE	M AND PACKING	INFORMATIO	N			
Is product exempt from DSCSA?	tion of manaractar	C1.	No	-	OLIV.	001722000000					,					
If yes, select exemption:				_	GCP:				i		Dimens	sions (US msr	nts)	Volume	Saleable #	
Other exemption - Write in:					GCI .				-1	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?			No		If yes, was or	riginal product			Item/Each:	0.00	3	3	6			
Is product sold by manufacturer's	exclusive distribu	tor?	Yes			irect from mfr?				0.63	3	3	ь	54.00	1	
Has FDA granted waiver/exceptio		oduct?	No		Provide sour	ce manufacturer f	for repa	ckaged product	Box/Carton/B	undle/						
If yes, attach documentation fro	m FDA.								Inner Pack:							
		0.7	TIN AND LUDGE PRODUCT IN	FORMATION.					Case:	8.25	13	9.75	6.75	855.56	12	
		G	TIN AND HIBCC PRODUCT IN	FORMATION					Pallet:			_				
Saleable Unit of Measure	99	aleable Quantity	HIBCC		GTI	N-14		Unit of Use GTIN-14	Pallet:							
X Item/Each		1	TIBOO			31722149051		5.11t of 030 OTHY-14								
Box/Carton/Bundle/Inner Pack	. 0000			31722143001			COST INFORMATION				WHOLESALER USE ONLY:					
x Case		12 1033			31722149058	1722149058										
Pallet								Regular Cost								
									Invoice Cost	(WAC) (\$)	\$40.00	Whsl. Code				
									II			Fineline Co	de:			
	-						-		As of date:							
									11							
			Attach copy of SAFETY DAT	A SHEET (SDS	S) or non hazar	rd letter PACKAGE	INSEPT	T LAREL AND PHOTO OF	PRODUCT PACK	AGING and BARCODE		-1				
*Please provide any additional inf		2	, audit copy of OAI ETT DAT	J. ILL I (JD	, oi noi nazai			nated Drop Ship Only.		Signature:						



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? No	SDS Hazard Classification						
b. CA Prop. 65 Carcinogen or Reproductive Toxicant? 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?	LEA Hazardous waste code.						
(if yes, answer a-e below and provide SDS) a. UN/Identification Number	REMS or REGISTRY RESTRICTIONS						
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: Phone: DEA #: NCPDP#: NCPDP#: NPI #:						
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments Registry: No						
ADD'L STORAGE INFORMATION	Registry: Registry Program Contact Name: Comments No Phone:						
Is the Product	RETURN INSTRUCTIONS						
Controlled Substance? Controlled Substance Code Controlled by State(s)? ARCOS Reportable? Schedule No. CLASS OF TRADE RESTRICTION: Controlled Substance Code Listed Chemical (List I or II) No If yes, indicate which: Is it a scheduled listed chemical product?: No	Contact tel. # if product received damaged: Is product returnable for credit: 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: Restricted to hospital, clinics, and physician offices only: No	Special regulations or returns requirements for this product in certain states? Yes						
Restricted from US territories? (explain in comments) Comments:	If so, which states? Other requirements? Comments? This product is classified as a schedule V controlled substance in Alabama, Kentucky, Michigan, North Dakota, Tennessee, Virginia, and West Virginia						
MISCELLANE	OUS NOTES and/or Image of Product Barcode:						



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

Version 2021

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?					