

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

Version 2021					Introduction 1	Type: Po:	st Launch Change		4 Final Version			Date:	10/20	/2023
		PRODUCT INFORMAT	ION						SPECIAL HANI	DLING AND STOR	AGE REQUIR	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA						ANDA	a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/AN	DA/BLA (drug); PMA/510(k)(m	ed device):	21495	7				Terr	perature Range	Controlled Room	- between 20	and 25 C (68	8° – 77° F)	
Medical Device Class, if applicable:														
	11-856-3719							Othe	er Temperature Range I	Requirement				
Proprietary Name (If Applicable) and		Gabapentin Tablets, USP 800 mg							(write in)					
J	31722-167-01	Unit of Use NDC:			UPC:	33172216701	7	Note	es					
UDI		CVX Code:			MVX Code:									
Description:	Gabapentin Tablets, USP 800	mg							is product to be shipped				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 Inform	nation: www.can	nberpharma.com						Nan		conons.	Soma Raju			
Address:	1031 Centennial Ave (and) 800				Address 2:			Nun	nber:		732-529-042	23		
	Piscataway			State:	NJ Zip: 08854			Group E-mail:			somaraju@heterousa.com			
	Customer Service			Email:	<u>customerservi</u>	ce@camber	pharma.com							
	1-866-827-3647			Fax:	732-562-8788				ons for product in any				No	
Product Therapeutic Classification	n: Anitconvuls	ant						Spe	cial returns requirement	ts for this product?			No	
		UCT INFORMATION		_	PRODUCT	ESCRIPTION	INFORMATION	d Chang and set (mit of colo) unviol: 10				No	
	ADDITIONAL PROD				PRODUCTI	DESCRIPTION	INFORMATION	• •	init of sale) upright?				No	
The product is?		Is the Product	Direct-Ship Only	/					tect product (unit of sa	ale) from light?			No	
a legend device? if yes, enter class #	No	Is the Product Orphan Drug Status	Neither		Size:	100 ct		e. Shelf life:	al shelf life at launch (if difforont).			24	Months Months
a product kit?	No	Orphan Drug Status				800 m	a	initi	ai sheir ille at launch (ir amerent):				wonths
if yes, list NDCs of	INU	FDA Approval Status			Strength:	800 11	g			ORDER INFORM	IATION			
component parts						Film o	oated tablets							
reverse numbered?	No				Dosage Forr	n:		Unit	of Sale		What is the	NDC selling	unit?	
co-licensed?	No	Allergens Present							x Bottle		1 Bottle of 10	00 Tablets		
latex-free?	Yes	Corn. Alco	hol, Wheat		Product Sha	Dval, I	piconvex		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?	Yes		inen, rineat		. roudor ond				Ampule					
correctional institution block?	No				Product Col	or: White			Glass		Minimum o	rder quantity	/?	Yes
opioid? Cannabinoid?	No	Country of Origin	USA			Debossed	with 'T' on the left side of the		Tube					
If Unit Dose, is item bar coded to un	No No	Country of Origin	USA		Product Imp	rint: bisect and	'3' on the right side of the bisect and bisect on other side.		Vial Liquid Sgl Vial Liquid Multi		If Yes how	many of whi	ch package	tvno?
hospital scanning?		Is this product covered up	nder the						Vial Powder Sql			Each	en package	type:
If Unit Dose, indicate NDC here:		Trade Agreements Act (T		es					Vial Power Multi			Inner/Carton	/Pack	
			1						Other: Write In			Case		
		FOR GENERIC DRUG PRO	DUCTS											
				Au	thorized Generic		Generic, other	PHARMACY ORDER / BILL UNIT						
	AB				section fields are not applicable			Rec. sell unit to customer? Rx billing unit to phare					acy:	
II. Generic Equivalent to What Bran	nd?: Neurontin	ablet						(Write-in, e.g. 1 Vial) Each						
	DRUG	SUPPLY CHAIN SECURITY ACT (D	OSCSA) INFORMA	TION				(write-in, e.g. 1 via	u)			Milliliter		
	5.00							1						
Does supplier meet DSCSA definit	tion of manufacturer?	Yes	GL	LN:	0331722000000				ITEM	AND PACKING I	NFORMATION	٩		
Is product exempt from DSCSA?		No												
If yes, select exemption:			GC	CP:					Weight Lbs.	Dimensi	ons (US msm	nts.)		Saleable #
Other exemption - Write in:									meigin Lus.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?		No			riginal product			Item/Each:	0.35	2.25	2.25	4.5	22.78	1
Is product sold by manufacturer's		Yes			irect from mfr?			D (0) (D)					-	
Has FDA granted waiver/exception If yes, attach documentation from		INO	Pr	ovide sour	ce manufacturer f	or repackaged	product	Box/Carton/Bundle Inner Pack:						
in yes, attach documentation non	III DA.							Case:						
		GTIN AND HIBCC PRODUCT IN	FORMATION						8.75	14	9.25	5.5	712.25	24
								Pallet:						
Saleable Unit of Measure	Saleable Qua	ntity HIBCC			N-14	Unit	of Use GTIN-14							
X Item/Each	1			003	31722167017									
Box/Carton/Bundle/Inner Pack				100	31722167014			(COST INFORMATION			WHOLESAL	ER USE ONL	in:
X Case Pallet	24	_		103	31722167014	-		Denular Cost			Vendor #:			
r anei								Regular Cost Invoice Cost (WAC	:) (\$)	\$20.00	Whsl. Code	#:		
									1 171	φ20.00	Fineline Co			
						1		As of date:						
						1					1			
Attach copy of SAFETY DATA SHEET (SDS) or non hazard letter, PACKAGE INSERT, LABEL AND PHOTO OF PRODUCT PACKAGING and BARCODE.														
*Please provide any additional info	ormation on page 2.				See new p. 3 for	Designated I	Drop Ship Only.	Sigi	nature:					

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Version 2021 For De	signated Drop Ship Only Products, Please Use Page 3								
MATERI	AL HAZARD CLASSIFICATION and TRANSPORTATION								
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?	Xo SDS Hazard Classification No X Organic Corrosive No Inorganic No Steroid/Androgen 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: No Identify NFPA Storage Level: No Is the product a NIOSH hazardous drug? No If yes, indicate which:								
c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	Hazardous Waste Identification EPA Hazardous Waste Code:								
Is this product regulated for shipment by IATA? (if yes, answer a-e below and provide SDS) a. UN/Identification Number b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	No REMS or REGISTRY RESTRICTIONS 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 Is this a reportable quantity?	No Med Guide Required Limited Distribution Requirement Comments / Details: (For example, iPledge program?) REMS: No No								
RQ Threshold: No 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) Special Permit; DOT-SP	REMS Program Manager Name: Phone: Supplier Manages REMS registry exclusively: Image: Comments								
Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Registry: No Registry Program Contact Name: Phone: Comments Phone:								
Is the Product Controlled Substance? No Controlled Substance Code Controlled by State(s)? Yes Listed Chemical (List I or II)	RETURN INSTRUCTIONS								
ARCOS Reportable? No If yes, indicate which:	No Contact tel. # if product received damaged: 1-866-827-3647 No Is product returnable for credit: Yes URL/Link to returns policy:								
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices	res contact - customerservice@camberpharma.com No Special regulations or returns requirements for this								
Restricted from US territories? (explain in comments)	No product in certain states? Yes No If so, which states? Other requirements? Comments?								
Comments:	This product is classified as a schedule V controlled substance in Alabama, Kentucky, Michigan, North Dakota, Tennessee, Virginia, and West Virginia								
MISCELLANEOUS NOTES and/or Image of Product Barcode:									



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Version 2021	FOR DESIGNATED DROP SHIP PRODUCT ONLY -	not a designated drop ship, do not complete.	
Order Method fo	r 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:	Fax Number: Fax Number: Phone No.: Site Address:	Purchase order daily receipt cut off time by supplier Cut off time: Shipping lead time of PO: Hours Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:	Days
1 3	Name:Phone:	-	_
Expedited Freight Charge	ges 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:			londay uesday /ednesday hursday riday
		Priority Overnight receipt available:	
Class	s of Trade Restriction:	PO Receipt Cut off time:	
No restriction: Select YES if sold to retail pha Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician of Restricted from US territories? (explain in cor Comments:		Saturday Overnight receipt available: PO Receipt Cut off time: PO Receipt Cut off time: Phone: Order receipt method: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:	
Other Data Info	rmation 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?	
Mi	scellaneous Notes:		
		ADDITIONAL INFORMATION	
		Is product order for scheduled patient procedure? Is product order for restocking purposes?	