

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

Version 2021						Introduction 1	Туре:	New Item	x	Final Version			Date:	6/6/2	2024
			PRODUCT INFORMA	TION			-			SPECIAL HAN	DLING AND STOR	AGE REQUIR	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc.				Applica	tion:	ANDA	a. Temperature – Indicate the USP temperature range for this product.								
	tion Number for NDA/ANDA/BLA (drug); PMA/510(k)(med device): 216800 Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)														
Medical Device Class, if applicable:															
DUNS:	11-856-3719							Other Temperature Range Requirement			Excursions permitted between 15° and 30°C (59°				
Proprietary Name (If Applicable) a		e: Buprop	ion Hydrochloride Extended		ts, USP (SR) 1					ite in)		and 86°F)			
Selling Unit NDC:	31722-066-05		Unit of Use NDC: CVX Code:			UPC: MVX Code:	331722	066051	Notes						
UDI						WIVA Code:									
Description:	Bupropion Hydrochlo	oride Extended-Rel	ease Tablets, USP (SR) 100	) mg						oduct to be shipped				No	
Active Ingredient(s): Bupropion hydrochloride, USP No															
Active ingredient(s). buildploin hydrocholide, open															
URL for Additional Product Inform	or Additional Product Information: www.camberpharma.com								Name: Soma Raju						
Address:	800 Centennial Ave, Suite 1			Address 2:				:		732-529-042	3				
City:	Piscataway				NJ		08854	Group E-mail: somaraju@heterousa.c			eterousa.cor	<u>n</u>			
Key Contact:		Customer Service Email:			customerservice	@camber	rpharma.com								
Phone Number:					732-562-8788			c. Special regulations for product in any states? No Special returns requirements for this product? No							
Product Therapeutic Classification	n: A	minoketone antide	pressant (NDRI)						Special	returns requirement	s for this product?			No	
ADDITIONAL PRODUCT INFORMATION PRODUCT DESCRIPTION INFORMATION d. Store product (unit of sale) upright? No															
The mediatie?	ADDITION			Direct-Ship C	)nly	I KOBOCT	SECONI				la) from l'atric				
The product is? a legend device?	N	0	Is the Product Is the Product	Neither	////			500 ct	e. Shelf life:	product (unit of sa	ile) from light?			No 24	Months
if yes, enter class #			Orphan Drug Status			Size:	c l	300 G		nelf life at launch (i	if different).			24	Months
a product kit?	N	lo	orphan Drug otatao			Of a second s	1	100 mg	initial of	ion nio at laanon (	an annon orney.				
if yes, list NDCs of			FDA Approval Status			Strength:		-			ORDER INFORM	IATION			
component parts						Dosage Form	m: F	Film coated tablet							
reverse numbered?		lo				, s			Unit of S			What is the		unit?	
co-licensed? latex-free?		lo 'es	Allergens Present				le la	Round, biconvex, bevel	x	Bottle Box/Carton		1 Bottle of 50	JU Tablets g. 1 Box of 10	) //iolo)	
preservative-free?		'es	C	)ye		Product Sha		edged		Ampule		(write-in, e.	y. I BUX UI II	J vidis)	
correctional institution block?		lo				Des des Col		Red		Glass		Minimum or	der quantity	?	Yes
opioid?	N	lo				Product Col				Tube				L	
Cannabinoid?	N	lo	Country of Origin	India		Product Imp		Imprinted with 'V1 48' on one side and plain on other side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	init dose for						5	side and plain on other side		Vial Liquid Multi				ch package t	ype?
hospital scanning? If Unit Dose, indicate NDC here:			Is this product covered u Trade Agreements Act (		No					Vial Powder Sgl Vial Powder Multi			Each Inner/Carton	/Dook	
Il Onit Dose, indicate NDC here.			Indde Agreenhents Act (		NO					Other: Write In			Case	Fack	
			FOR GENERIC DRUG PR	ODUCTS									ouso		
					Au	uthorized Generic		orized Generic, other		PH	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB1						section	fields are not applicable	Rec. sell unit to custor	ner?		Rx billing u	nit to pharma	acy:	
II. Generic Equivalent to What Brand?: Wellbutrin SR									Each						
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION     Gram															
		DRUG SUPPL	Y CHAIN SECURITY ACT (	DSCSA) INFOR	MATION				-				Milliliter		
Does supplier meet DSCSA definit	tion of manufacturer	2	Yes	_	GLN:	0331722498975				ITEN	I AND PACKING I	NFORMATION	J		
Is product exempt from DSCSA?			No	-	02.11	0001122100010									
If yes, select exemption:					GCP:						Dimensi	ons (US msm	its.)	Volume	Saleable #
Other exemption - Write in:										Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?	_		No		If yes, was o	riginal product pur	chased		Item/Each:	0.55	2.87	2.87	5	41.18	1
Is product sold by manufacturer's			Yes	_	direct from n					0.00	2.07	2.07		41.10	
Has FDA granted waiver/exception		uct?	No		Provide sour	ce manufacturer fo	or repack	aged product	Box/Carton/Bundle/						
If yes, attach documentation from	n FDA.								Inner Pack: Case:						
		GTI	N AND HIBCC PRODUCT I	NFORMATION					Case.	7.1	12	9.18	5.75	633.42	12
									Pallet:						
Saleable Unit of Measure	Sale	eable Quantity	HIBCC			IN-14		Unit of Use GTIN-14							
X Item/Each		1			003	31722066051	_								
Box/Carton/Bundle/Inner Pack		10				24700000000	-		COS	T INFORMATION			WHOLESALI	ER USE ONL	Y:
X Case		12			303	31722066052	-		Regular Cost			Vendor #:			
					-		-		Invoice Cost (WAC) (\$		\$100.00	Whsl. Code	#:		
											\$100.00	Fineline Cod			
									As of date:	10/16/2023					
												1			
μ									L <u>I</u>			ļ			
			Attach copy of SAFETY DA	ATA SHEET (SD	S) or non haza			, LABEL AND PHOTO OF P							
*Please provide any additional info	ormation on page 2.					See new p. 3 for	r Designa	ated Drop Ship Only.	Signatu	re:					

## HDA🔾

## **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? 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? 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 Is this product regulated for shipment by DOT? (if yes, answer a-e below and provide SDS) a. UN/Identification Number No No No No No No No N	Does the product have an Aerosol class? If yes, identify NFPA Storage Level:       No         NFPA Storage Level:       Is the product a NIOSH hazardous drug?         Is the product a NIOSH hazardous drug?       No         If yes, indicate which:       If yes, indicate which:							
a. On/definition for holder     b. Proper Shipping Name     c. DOT Hazard Class     d. Packing Group     e. Inhalation Hazard?     Is this product regulated for shipment by IATA?     No	Hazardous Waste Identification EPA Hazardous Waste Code: Waste Code:							
(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?	REMS or REGISTRY RESTRICTIONS         Is there a REMS on this product?       No         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) Special Permit; DOT-SP	REMS:     No       REMS Program Manager Name:     Phone:       Supplier Manages REMS registry exclusively:     Phone:       Wholesale distributor support:     DEA #:       Provider Name:     DEA #:       Site Enrollment Number assigned     NCPDP#:       by Supplier:     NPI #:							
Special Provision (listed in Column 7 of 49 CFR 172.101); SP#ADD'L STORAGE INFORMATION	No       Registry Program Contact Name:       Comments							
Is the Product Controlled Substance? Controlled Substance? No Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which: Schedule No. Is it a scheduled listed chemical product?: No	RETURN INSTRUCTIONS         Contact tel. # if product received damaged:       1-866-827-3647         Is product returnable for credit:       Yes							
CLASS OF TRADE RESTRICTION: No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Yes	URL/Link to returns policy: contact - customerservice@camberpharma.com							
No     No       Restricted to retail pharmacy only:     No       Restricted to hospital, clinics, and physician offices only:     No       Restricted from US territories? (explain in comments)     No       Comments:     No	Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?							
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         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:       Image: Comparison of the co
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:         Order receipt method:       Phone:         Fax:       EDI:         Overnight Fees apply:       Other fees apply:
Other Data Information Required to Process PO:	Return Instructions
Patient Procedure Date:	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?