

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

Version 2021					Introduction T	ype: Post Launch Change	x	Final Version			Date:	11/19	9/2024
		PRODUCT INFORMA	TION					SPECIAL HAN	DLING AND STOP	RAGE REQUIR	EMENTS*	m	
Company Name:	Camber Pharmaceuticals. Inc				Applicat	ion: ANDA	a. Temperature – Indi	cate the USP tempe	erature range for t	his product.			
Application Number for NDA/AN			202	682		· .		ature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicat		· ·											
DUNS:	11-856-3719						Other T	emperature Range I	Requirement				
Proprietary Name (If Applicable) a	and Established Name:	Atomoxetine Hydrochloride Capsu					(v	rite in)					
Selling Unit NDC:	31722-715-30	Unit of Use NDC:		31722-715-30		331722715300	Notes						
UDI		CVX Code:			MVX Code:								
Description:	Atomoxetine Hydrochloride C	apsules, USP 18 mg					Is this p	roduct to be shipped	d to customers on i	ce?		No	1
Is this product to be shipped to customers on dry ice? No													
Active Ingredient(s):	Atomoxeti	ne hydrochloride, USP											
b. Contact for temperature excursion questions:													
URL for Additional Product Inform Address:		erpharma.com			Address 2:		Name:			Soma Raju			
City:	800 Centennial Ave, Suite 1 Piscataway			State:	NJ	Zip: 08854	Group			732-529-042 somaraju@h		•	
Key Contact:	Customer Service			Email:		camberpharma.com	Group	E-IIIdii.		somarajuen	eterousa.con	<u>u</u>	
Phone Number:	1-866-827-3647			Fax:	732-562-8788	<u>soundorphamatoonn</u>	c. Special regulations	for product in any	states?			No	1
Product Therapeutic Classification		orepinephrine reuptake inhibitor (SN	RI)					returns requirement				No	
			,				opoola	rotanio roquironion					1
	ADDITIONAL PRO	DUCT INFORMATION			PRODUCT	DESCRIPTION INFORMATION	d. Store product (unit	of sale) upright?				No	1
The product is?		Is the Product	Direct-Ship Or	nlv					ale) from light?			No	1
The product is? a legend device?	No	Is the Product	Unit of Use	,		30 ct	e. Shelf life:	product (unit of sa	ae, nom nynt?			24	Months
if yes, enter class #		Orphan Drug Status	5		Size:	00 01		helf life at launch (	if different):			24	Months
a product kit?	No					18 mg							1
if yes, list NDCs of		FDA Approval Status			Strength:				ORDER INFORM	IATION			
component parts					Dosage Form	Hard gelatin capsule							
reverse numbered?	No				Dosageron		Unit of			What is the		unit?	
co-licensed?	No	Allergens Present					X	Bottle		1 Bottle of 30			
latex-free?	Yes	Animal	Products		Product Sha	capsule		Box/Carton		(Write-in, e.	g. 1 Box of 10	) Vials)	
preservative-free?	Yes					Velley, een and uhite		Ampule		Minimum		•	Vaa
correctional institution block? opioid?	No				Product Cold	Yellow cap and white opaque body		Glass Tube		Minimum or	der quantity	?	Yes
Cannabinoid?	No	Country of Origin	India			Imprinted with 'l' on cap		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u		country of origin	India		Product Imp	and '106' on body		Vial Liquid Multi		If Yes, how	manv of whi	ch package t	type?
hospital scanning?		Is this product covered	under the					Vial Powder Sgl			Each		.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
If Unit Dose, indicate NDC here:		Trade Agreements Act (		No				Vial Powder Multi			Inner/Carton	/Pack	
			5					Other: Write In			Case		
		FOR GENERIC DRUG PR	ODUCTS										
								5					
				Au	thorized Generic	*If Authorized Generic, other		PH	IARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB					section fields are not applicable	Rec. sell unit to custo	mer?		Rx billing u	it to pharma	acy:	
II. Generic Equivalent to What Bra	nd?: Strattera							Each					
							(Write-in, e.g. 1 Vial)				Gram		
	DRU	G SUPPLY CHAIN SECURITY ACT	(DSCSA) INFORI	MATION							Milliliter		
Doos supplier most DSCSA definit	tion of manufacturar?	Vac	_	CLN.	0331722498975			ITEN	I AND PACKING I				
Does supplier meet DSCSA definit Is product exempt from DSCSA?		Yes No		GLN:	03317224909/5			11 EN	HAND FACKING I				
		110							Dime		ta )		0-1 "
If yes, select exemption:				GCP:				Weight Lbs.		ions (US msm Width	,	Volume (Cube)	Saleable # Pieces
Other exemption - Write in: Is product repackaged?		No		If yos was as	iginal product pure	hased	Item/Each:		Depth	Width	Height		
				direct from m			nem/Lacit.	0.07	1.55	1.55	2.55	6.13	1
Is product sold by manufacturer's	exclusive distributor?	Yes				- reveal and an dust							
Is product sold by manufacturer's Has FDA granted waiver/exception		Yes		Provide source	ce manufacturer fo	r repackaged product	Box/Carton/Bundle/						
Is product sold by manufacturer's Has FDA granted waiver/exception If yes, attach documentation from	n/exemption for product?			Provide source	ce manufacturer fo	r repackaged product	Box/Carton/Bundle/ Inner Pack:						0.4
Has FDA granted waiver/exception	n/exemption for product?			Provide source	ce manufacturer fo	r repackaged product		2	9.75	6.75	4	263.25	
Has FDA granted waiver/exception	n/exemption for product?			Provide source	ce manufacturer to	r repackaged product	Inner Pack: Case:	2	9.75	6.75	4	263.25	24
Has FDA granted waiver/exception If yes, attach documentation from	n/exemption for product? m FDA.	No GTIN AND HIBCC PRODUCT I					Inner Pack:	2	9.75	6.75	4	263.25	24
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure	n/exemption for product? m FDA. Saleable Qua	No GTIN AND HIBCC PRODUCT I		GTII	N-14	Unit of Use GTIN-14	Inner Pack: Case:	2	9.75	6.75	4	263.25	24
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure	n/exemption for product? m FDA.	No GTIN AND HIBCC PRODUCT I		GTII			Inner Pack: Case: Pallet:		9.75				
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X Item/Each Box/Carton/Bundle/Inner Pack	n/exemption for product? m FDA. Saleable Que	No GTIN AND HIBCC PRODUCT I		GTII 003:	N-14 31722715300	Unit of Use GTIN-14	Inner Pack: Case: Pallet:	2 ST INFORMATION	9.75			263.25 ER USE ONL	
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure	n/exemption for product? m FDA. Saleable Qua	No GTIN AND HIBCC PRODUCT I		GTII 003:	N-14	Unit of Use GTIN-14	Inner Pack: Case: Pallet:		9.75				
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X Item/Each Box/Cartor/Bundle/Inner Pack Case	n/exemption for product? m FDA. Saleable Que	No GTIN AND HIBCC PRODUCT I		GTII 003:	N-14 31722715300	Unit of Use GTIN-14	Inner Pack: Case: Pallet: Cox Regular Cost	ST INFORMATION		Vendor #:	WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X Item/Each Box/Cartor/Bundle/Inner Pack Case	n/exemption for product? m FDA. Saleable Que	No GTIN AND HIBCC PRODUCT I		GTII 003:	N-14 31722715300	Unit of Use GTIN-14	Inner Pack: Case: Pallet:	ST INFORMATION			WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X Item/Each Box/Cartor/Bundle/Inner Pack Case	n/exemption for product? m FDA. Saleable Que	No GTIN AND HIBCC PRODUCT I		GTII 003:	N-14 31722715300	Unit of Use GTIN-14	Inner Pack: Case: Pallet: Cox Regular Cost	ST INFORMATION		Vendor #: Whsl. Code	WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X Item/Each Box/Cartor/Bundle/Inner Pack Case	n/exemption for product? m FDA. Saleable Que	No GTIN AND HIBCC PRODUCT I		GTII 003:	N-14 31722715300	Unit of Use GTIN-14	Inner Pack: Case: Pallet: CO Regular Cost Invoice Cost (WAC) (\$	ST INFORMATION		Vendor #: Whsl. Code	WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X Item/Each Box/Cartor/Bundle/Inner Pack Case	n/exemption for product? m FDA. Saleable Que	No       GTIN AND HIBCC PRODUCT I       Intity       HIBCC	NFORMATION	GTII 003: 203:	N-14 31722715300 31722715304	Unit of Use GTIN-14 00331722715300	Inner Pack: Case: Pallet: Regular Cost Invoice Cost (WAC) (\$ As of date:	ST INFORMATION (1)		Vendor #: Whsl. Code	WHOLESALI		
Has FDA granted waiver/exception If yes, attach documentation from Saleable Unit of Measure X Item/Each Box/Cartor/Bundle/Inner Pack X Case	n/exemption for product? m FDA.	No       GTIN AND HIBCC PRODUCT I       Intity       HIBCC	NFORMATION	GTII 003: 203:	N-14 31722715300 31722715304 rd letter, PACKAGE	Unit of Use GTIN-14	Inner Pack: Case: Pallet: Regular Cost Invoice Cost (WAC) (\$ As of date:	ST INFORMATION		Vendor #: Whsl. Code	WHOLESALI		

## HDA🔾

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

Version 2021 For Designa	ted Drop Ship Only Products, Please Use Page 3					
MATERIAL HA	ZARD 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					
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					
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?