

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

Version 2021					Introduction	Туре:	New Item		x Final Ve	rsion			Date:	3/20/	2024
		PRODUCT INFORMA	TION						SPEC	IAL HANI	DLING AND STOR	AGE REQUIR	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc.				Applica	Application: ANDA			a. Temperature – Indicate the USP temperature range for this product.							
Application Number for NDA/ANI	DA/BLA (drug); PMA/510(k)(me	d device):	204792						Temperature Ran	ge .	Controlled Room	- between 20	and 25 C (68	° – 77° F)	
Medical Device Class, if applicab															
DUNS:	11-856-3719								Other Temperatur	e Range R	Requirement				
Proprietary Name (If Applicable) an		Fesoterodine Fumarate Extended		000.00					(write in)						
Selling Unit NDC: UDI	31722-033-30	Unit of Use NDC: CVX Code:	31722-	-033-30	UPC: MVX Code:	33172203	3305		Notes						
					intx oouc.										
Description:	Fesoterodine Fumarate Extende	ed-Release Tablets 4 mg							Is this product to b					No No	
Active Ingredient(s): Is this product to be shipped to customers on dry ice? No															
b. Contact for temperature excursion questions:															
URL for Additional Product Inform									Name:			Soma Raju			
Address:	1031 Centennial Ave (and) 800	Centennial Ave, Suite 1			Address 2:				Number:			732-529-042			
City:	Piscataway Customer Service			tate: mail:	NJ customerservice	Zip: 0			Group E-mail:			somaraju@h	eterousa.cor	<u>1</u>	
Key Contact: Phone Number:	1-866-827-3647			ax:	732-562-8788	e camperpri	lama.com	c. Special reg	ulations for produ	ct in any	states?			No	
Product Therapeutic Classification		nic							Special returns re	-				No	
	-								opoolariotanio ro	quironnonna					
	ADDITIONAL PRODU				PRODUCT	DESCRIPTI	ION INFORMATION	d. Store produ	uct (unit of sale) u	pright?				No	
The product is?		Is the Product	Direct-Ship Only					1	Protect product (le) from liaht?			No	
a legend device?	No	Is the Product	Unit of Use	_	Circ.	30	ct	e. Shelf life:			,			24	Months
if yes, enter class #		Orphan Drug Status			Size:				Initial shelf life at	launch (i	f different):				Months
a product kit?	No				Strength:	4 m	ng				-				
if yes, list NDCs of		FDA Approval Status			onongan						ORDER INFORM	IATION			
component parts reverse numbered?					Dosage For	m: Filr	m coated tablet		Unit of Sale			What is the		uni#2	
co-licensed?	No	Allergens Present							x Bottle			1 Bottle of 30		unit	
latex-free?	Yes					Ov	al, biconvex		Box/Car	ton			g. 1 Box of 10) Vials)	
preservative-free?	Yes		Soy		Product Sha	ape:	,		Ampule			(J		
correctional institution block?	No				Product Co	Lig	ht blue		Glass			Minimum or	der quantity	?	Yes
opioid?	No			_	Troduct CO				Tube						
Cannabinoid?	No	Country of Origin	India		Product Imp		bossed with 'H' on one		Vial Liqu			W. V			
If Unit Dose, is item bar coded to un hospital scanning?	nit dose for	Is this product covered u	inder the			sid	le and 'F6' on the other		Vial Liqu Vial Pov				many of whit Each	ch package t	sype?
If Unit Dose, indicate NDC here:		Trade Agreements Act (0.d			Vial Pov				Inner/Carton	/Pack	
		······	110						Other: V				Case	aon	
		FOR GENERIC DRUG PR	ODUCTS												
				Au	thorized Generic		ized Generic, other			PH.	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating: AB section fields are not applicable							elds are not applicable	Rec. sell unit to customer? Rx billing unit to pharmacy:							
II. Generic Equivalent to What Brand?: Toviaz								Each							
	DRUC	SUPPLY CHAIN SECURITY ACT		M				(Write-in, e.g.	1 Vial)				Gram Milliliter		
	DRUG	SUPPLY CHAIN SECORITY ACT	DSCSA) INFORMATIO	N				-					Millinter		
Does supplier meet DSCSA definit	tion of manufacturer?	Yes	GLN:		0331722000000			-		ITEM	AND PACKING I	NFORMATION	N		
Is product exempt from DSCSA?		No													
If yes, select exemption:			GCP:					1			Dimensi	ons (US msm	nts.)	Volume	Saleable #
Other exemption - Write in:									Weig	ht Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?		No			riginal product pu	rchased		Item/Each:).1	1.56	1.56	3.13	7.62	1
Is product sold by manufacturer's		Yes		from m											
Has FDA granted waiver/exception If yes, attach documentation from		No	Provid	e sour	ce manufacturer f	or repackag	jea product	Box/Carton/Bo Inner Pack:	undle/						
in yes, attach documentation from								Case:				0			
		GTIN AND HIBCC PRODUCT I	NFORMATION						2	2.8	9.75	6.75	4	263.25	24
								Pallet:							
Saleable Unit of Measure	Saleable Quan	tity HIBCC			N-14		Init of Use GTIN-14								
X Item/Each	1			003	31722033305	0	0331722033305								
Box/Carton/Bundle/Inner Pack	24			202	31722033309	_			COST INFOR	MATION			WHOLESALI	ER USE ONL	r:
X Case	24			203.	31722033309	-		Regular Cost				Vendor #:			
						-		Invoice Cost (WAC) (\$)		\$40.00	Whsl. Code	#:		
						-					÷.0.00	Fineline Co			
								As of date:	3/14/202	24					
								1							
μ								1				1			
		Attach copy of SAFETY D	ATA SHEET (SDS) or no	on haza						DE.					
*Please provide any additional information on page 2. See new p. 3 for Designated Drop Ship Only. Signature:															

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

·	nated Drop Ship Only Products, Please Use Page 3
MATERIAL H	HAZARD CLASSIFICATION and TRANSPORTATION
Is this product (check all that apply): a. Cytotoxic? No b. CA Prop. 65 Carcinogen or Reproductive Toxicant? Is the product a CA Prop 65 carcinogen? No Is the product a CA Prop 65 reproductive toxicant? No Does the product label bear a CA Prop 65 warning? C. Contact Hazard? (If yes, attach SDS with special instructions.) e. Does the product contain DEHP? No	x Organic Corrosive Inorganic Oxidizer Steroid/Androgen Contact Hazard Does the product have an Aerosol class? If yes, identify NFPA Storage Level: No NFPA Storage Level: Image: Contact Hazard
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 c. DOT Hazard Class	Is the product a NIOSH hazardous drug? No If yes, indicate which: Hazardous Waste Identification
d. Packing Group 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) a. UN/Identification Number b. Proper Shipping Name	REMS or REGISTRY RESTRICTIONS Is there a REMS on this product? No
c. DOT Hazard Class d. Packing Group e. Inhalation Hazard? Is the product restricted for air shipment? If so, indicate restriction: Passenger	If Yes, is it managed with a pharmacy registry? Website URL: Med Guide Required Limited Distribution Requirement
Cargo Passenger & Cargo Is this a reportable quantity? No RQ Threshold:	REMS: No REMS Program Manager Name: Phone:
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)	Supplier Manages REMS registry exclusively:
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101);	Comments
ADD'L STORAGE INFORMATION	Registry: No Registry Program Contact Name: Phone: Comments
Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS
Controlled by State(s)? No Listed Chemical (List I or II) No ARCOS Reportable? No If yes, indicate which: Image: Control of the state of	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: No Restricted to hospital, clinics, and physician offices only: No Restricted from US territories? (explain in comments) No	Special regulations or returns requirements for this product in certain states? No If so, which states? Other requirements? Comments?
Comments:	
MISCELLAN	IEOUS NOTES and/or Image of Product Barcode:



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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?