

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

Version 2021						Introduction 1	Туре:	New Item	x	Final Version			Date:	6/24/	/2024
			PRODUCT INFORMA	TION			-	-		SPECIAL HAN	DLING AND STOP	RAGE REQUI	REMENTS*		
Company Name:	Camber Pharmac	ceuticals. Inc.				Applica	tion:	ANDA	a. Temperature – Ind	icate the USP tempe	erature range for t	his product.			
Application Number for NDA/ANI			evice):	215	5742					rature Range	Controlled Room		and 25 C (68	° – 77° F)	
Medical Device Class, if applicab			,							J. J					
DUNS:	11-856-3719								Other	Femperature Range I	Requirement	Excursions p	permitted betw	ween 15°C to	
Proprietary Name (If Applicable) and	nd Established Na	ame: Zile	euton Extended-Release Table						()	write in)		30°C (59°F t	o 86°F)		
Selling Unit NDC:	31722-044-12		Unit of Use NDC		31722-044-12		331722	2044127	Notes						
UDI			CVX Code:			MVX Code:									
Description:	Zileuton Extended	d-Release Tablets	600 mg						Is this	product to be shipped	d to customers on i	ce?		No	
Is this product to be shipped to customers on dry ice? No															
Active Ingredient(s): Zileuton, USP															
UDL (as Additional Desident Inform									b. Contact for temper		estions:	Soma Raju			
URL for Additional Product Inform Address:		www.camberpha	arma.com		1	Address 2:			Name:			732-529-042	22		
City:	800 Centennial Ave, Suite 1 Piscataway State:				NJ Zip: 08854			Number: Group E-mail:			eterousa.cor	n			
Key Contact:	Customer Service Email:				omerservice@camberpharma.com				Somaraja en	101010434.001	<u>n</u>				
Phone Number:	1-866-827-3647				732-562-8788				s for product in any	states?			No	1	
Product Therapeutic Classification	ו:	Leukotriene syn	thesis inhibitor							I returns requirement				No	
										1					
	ADDITI	IONAL PRODUCT				PRODUCT	DESCRI	IPTION INFORMATION	d. Store product (uni	t of sale) upright?				No]
The product is?			Is the Product	Direct-Ship C	Dnly				Protec	t product (unit of sa	le) from liaht?			No	1
a legend device?		No	Is the Product	Unit of Use		Sizo	[120 ct	e. Shelf life:		,			24	Months
if yes, enter class #			Orphan Drug Status			Size:				shelf life at launch (if different):				Months
a product kit?		No				Strength:		600 mg							
if yes, list NDCs of			FDA Approval Status			cuongan					ORDER INFORM	IATION			
component parts						Dosage Forr	m:	Bi-layer, film-coated tablet	11-21-21	0-1-		What is the	NDC selling		
reverse numbered?		No	Allergens Present				l.		Unit of	Bottle		1 Bottle of 1		unit?	
co-licensed? latex-free?		No Yes	_					Oblong, biconvex		Box/Carton			g. 1 Box of 1	() (Vials)	
preservative-free?		Yes	c	orn		Product Sha	ape:	Obiolity, bioditiex		Ampule		(write iii, e.	g. 1 Dox 01 1	0 100)	
correctional institution block?		No				Des test Oct		Pink to red IR layer on one side and		Glass		Minimum o	der quantity	?	Yes
opioid?		No				Product Cole		white to off white ER layer on the other side		Tube					
Cannabinoid?		No	Country of Origin	India		Product Imp	nint.	Debossed '66' on pink to red side and 'V' on the white to off white side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	nit dose for					ouuot imp		V on the white to on white side		Vial Liquid Multi				ch package t	type?
hospital scanning?			Is this product covered Trade Agreements Act (N					Vial Powder Sgl		12	Each	(De ele	
If Unit Dose, indicate NDC here:			Trade Agreements Act (TAA)?	No					Vial Powder Multi Other: Write In			Inner/Carton Case	Ласк	
			FOR GENERIC DRUG PF	ODUCTS									Case		
			TOK GENERIC DROG FI	000013					-						
					Au	thorized Generic	*If Auth	horized Generic, other	-	PH	ARMACY ORDER	/ BILL UNIT			
I. Orange Book Rating:	AB							n fields are not applicable	Rec. sell unit to cust	omer?		Ry billing u	nit to pharm:	acv:	
II. Generic Equivalent to What Brand?: Zyflo CR							Rec. sell unit to customer? Rx billing unit to pharmacy:								
							(Write-in, e.g. 1 Vial)		-		Gram				
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFORMATION Milliliter															
			No							1751					
Does supplier meet DSCSA definit Is product exempt from DSCSA?	tion of manufactur	rer?	Yes		GLN:	0331722498975				IIEN	I AND PACKING II	NFORMATIO	N		
											D				
If yes, select exemption:					GCP:					Weight Lbs.		ions (US msn	,	Volume	Saleable #
Other exemption - Write in: Is product repackaged?			No		If yoe was	iginal product pur	chacod		Item/Each:	-	Depth	Width	Height	(Cube)	Pieces
Is product sold by manufacturer's	exclusive distribu	utor?	Yes	-	direct from m		chaseu		nem/Each.	0.35	2.6	2.6	4.21	28.46	1
Has FDA granted waiver/exception			No	-		ce manufacturer fo	or repacl	kaged product	Box/Carton/Bundle/						
If yes, attach documentation from		L							Inner Pack:						
									Case:	4.7	11	8.75	6	577.50	12
		(GTIN AND HIBCC PRODUCT	NFORMATION								0.10		011.00	
Saleable Unit of Measure	-							Unit of the OTTO A	Pallet:						
	S	Saleable Quantity	HIBCC			N-14		Unit of Use GTIN-14 00331722044127							
X Item/Each Box/Carton/Bundle/Inner Pack		1 00331722044127 00331722044127						00001122044121		ST INFORMATION			WHOLESAL	ER USE ONL	Y:
X Case		12 20331722044121					COSTINFORMATION				WHOLESALER USE ONLY:				
Pallet					200				Regular Cost			Vendor #:			
	1						1		Invoice Cost (WAC) (\$)	\$600.00	Whsl. Code	#:		
]											Fineline Co			
									As of date:	3/7/2023		ļ			
]											1			
μ									<u> </u>			ļ			
			Attach copy of SAFETY D	ATA SHEET (SD	OS) or non haza			T, LABEL AND PHOTO OF P							
*Please provide any additional info	ormation on page	2.				See new p. 3 for	r Design	ated Drop Ship Only.	Signat	ure:					

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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?