

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

Version 2021					Introduction Type	: New Item		x Final Version			Date:	6/24/	1/2024
			PRODUCT INFORMA	TION				SPECIAL HAN	DLING AND STOR	RAGE REQUI	REMENTS*		
Company Name: Camber Pharmaceuticals, Inc. Application: ANDA							a. Temperature – Indicate the USP temperature range for this product.						
Application Number for NDA/AN	IDA/BLA (drug); PM	A/510(k)(med device	ce):	212475		<u> </u>		Temperature Range	Controlled Room		and 25 C (68	3° – 77° F)	
Medical Device Class, if applical				·			1	· -					
DUNS:	11-856-3719							Other Temperature Range F	Requirement				
Proprietary Name (If Applicable) a	and Established Nar	me: Zafirlu	ikast Tablets 10 mg					(write in)					
Selling Unit NDC:	31722-007-60		Unit of Use NDC:	31722-007-60		1722007603		Notes					
UDI			CVX Code:		MVX Code:								
Description:	Zafirlukast Tablets	10 mg					Ī	Is this product to be shipped	to customers on i	ice?		No	1
-								Is this product to be shipped				No	1
Active Ingredient(s):		Zafirlukast											
								r temperature excursion que	estions:				
URL for Additional Product Inforn		www.camberpharm	a.com					Name:		Soma Raju			
Address:	800 Centennial Ave	e, Suite 1		Ctata	Address 2:	00054		Number:		732-529-042			
City:	Piscataway Customer Service			State: Email:	NJ Zi customerservice@car	ip: 08854	Group E-mail: somaraju@heterousa.com						
Key Contact: Phone Number:	1-866-827-3647			Fax:	732-562-8788	mberpriarma.com	a Special rea	julations for product in any	ctotoc?			No	٦
Product Therapeutic Classificatio		Synthotic coloctive r	peptide leukotriene receptor a		732-302-0700		c. Special reg	Special returns requirement				No	-
Product Therapeutic Classificatio	on:	Synthetic, Selective p	peptide leukotherie receptor a	magoriisi (LTRA)				Special returns requirement	s for this product?			INO	_
	ADDITIO	NAL PRODUCT IN	FORMATION		PRODUCT DES	CRIPTION INFORMATION	d Store prod	uct (unit of sale) upright?				No	٦
	ADDITIO	MAET NODOOT IN		Discoul Obis Oak	TRODUCT DEC	SKII TISK IKI SKIIIATISK	u. Store prou						4
The product is?		NI.	Is the Product	Direct-Ship Only Unit of Use		00 -1	. 01-1/17	Protect product (unit of sa	le) from light?			No 04	
a legend device? if yes, enter class #		No	Is the Product Orphan Drug Status	Offit of Ose	Size:	60 ct	e. Shelf life:	Initial shelf life at launch (if different).			24	Months Months
a product kit?		No	Orphan Drug Status			10 mg		initial shell life at launch (ir different):				Wonths
if yes, list NDCs of		140	FDA Approval Status		Strength:	To mg			ORDER INFOR	MATION			
component parts						Film coated tablet							
reverse numbered?		No			Dosage Form:			Unit of Sale		What is the	NDC selling	unit?	
co-licensed?		No	Allergens Present					x Bottle		1 Bottle of 6	0 Tablets		
latex-free?		Yes	La	ctose	Product Shape:	Round, biconvex		Box/Carton		(Write-in, e.	g. 1 Box of 1	0 Vials)	
preservative-free?		Yes	Lu	Close	i roduct onapc.			Ampule					
correctional institution block?		No			Product Color:	White to off-white		Glass		Minimum o	rder quantity	/?	Yes
opioid?		No						Tube					
Cannabinoid?		No	Country of Origin	India	Product Imprint:	Debossed with 'V' on one side and '16' on the other side		Vial Liquid Sgl					
If Unit Dose, is item bar coded to u	unit dose for		In this was dead a second of	and and the	·	and 16 on the other side		Vial Liquid Multi Vial Powder Sql			many of wh	ich package i	type?
hospital scanning? If Unit Dose, indicate NDC here:			Is this product covered to Trade Agreements Act (Vial Powder Sgi Vial Powder Multi		24	Inner/Cartor	/Pook	
Il Ollit Dose, ilidicate NDC liele.			Trade Agreements Act (IAA): NO				Other: Write In			Case	I/Fack	
			FOR GENERIC DRUG PR	PODLICTS			1	Other: Write in			Joase		
			TOR GENERIC DROGT	.000013									
				A	uthorized Generic *If	Authorized Generic, other		PH	ARMACY ORDER	/ BILL UNIT			
I Oranga Baak Batings	AB					ction fields are not applicable	Pac sall unit	to customer?					,
I. Orange Book Rating: II. Generic Equivalent to What Bra		Accolate					Nec. sen unit	to customer:	1	KX billing u	nit to pharm Each	acy:	
II. Generic Equivalent to what Bra	anur.	Accorate					(Write-in, e.g	1 Vial)	J		Gram		
		DRUG SUPPL	LY CHAIN SECURITY ACT	(DSCSA) INFORMATION			(**************************************	· · · · · · · · ·			Milliliter		
				, ,									
Does supplier meet DSCSA defini	ition of manufacture	er?	Yes	GLN:	0331722498975			ITEN	I AND PACKING I	NFORMATIO	N		
Is product exempt from DSCSA?			No					·					
If yes, select exemption:				GCP:			1		Dimens	ions (US msn	nts.)	Volume	Saleable #
Other exemption - Write in:							1	Weight Lbs.	Depth	Width	Height	(Cube)	Pieces
Is product repackaged?			No	If yes, was o	riginal product purchas	sed	Item/Each:	0.06	1.5	1.5	3.5	7.88	1
Is product sold by manufacturer's			Yes	direct from r					1.5	1.5	3.3	7.00	
Has FDA granted waiver/exceptio		oduct?	No	Provide sour	ce manufacturer for re	packaged product	Box/Carton/E	Bundle/					
If yes, attach documentation from	m FDA.						Inner Pack:						
		CTI	IN AND HIBCC PRODUCT I	NEODMATION			Case:	2.1	9.5	6.5	4.5	277.88	24
		GII	IN AND RIBCC PRODUCT I	NFORMATION			Pallet:						
Saleable Unit of Measure	99	aleable Quantity	HIBCC	GT	IN-14	Unit of Use GTIN-14	Fallet.						
X Item/Each	36	1	TIIDOO		331722007603	00331722007603							
Box/Carton/Bundle/Inner Pack				000				COST INFORMATION			WHOLESAL	ER USE ONL	LY:
X Case		24		203	331722007607								
Pallet							Regular Cost			Vendor #:			
							Invoice Cost	(WAC) (\$)	\$73.74	Whsl. Code	#:		
										Fineline Co	de:		
							As of date:	11/9/2020					
							1 1						
 					510//165		DODUGE DATE						
*Please provide any additional inf			Attach copy of SAFETY D.	ATA SHEET (SDS) or non haza		SERT, LABEL AND PHOTO OF F	PRODUCT PACK	AGING and BARCODE. Signature:		-			



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? No	SDS Hazard Classification						
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? No 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?	Does the product have an Aerosol class? If yes, identify NFPA Storage Level: NFPA Storage Level:						
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	Is the product a NIOSH hazardous drug? If yes, indicate which:						
c. DOT Hazard Class d. Packing Group	Hazardous Waste Identification						
e. Inhalation Hazard? Is this product regulated for shipment by IATA? No	EPA Hazardous Waste Code: Waste Characteristics						
(if yes, answer a-e below and provide SDS) a. UN/Identification Number	REMS or REGISTRY RESTRICTIONS						
b. Proper Shipping Name c. DOT Hazard Class d. Packing Group e. Inhalation Hazard?	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	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? Yes 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)	REMS: REMS Program Manager Name: Supplier Manages REMS registry exclusively: Wholesale distributor support: Provider Name: Site Enrollment Number assigned by Supplier: No Phone: DEA #: NCPDP#: NCPDP#: NPI #:						
Special Permit; DOT-SP Special Provision (listed in Column 7 of 49 CFR 172.101); SP#	Comments						
ADD'L STORAGE INFORMATION	Registry: Registry Program Contact Name: Comments No Phone:						
Is the Product							
Controlled Substance Code Controlled by State(s)? ARCOS Reportable? Schedule No. No Listed Chemical (List I or II) No If yes, indicate which: Is it a scheduled listed chemical product?: No CLASS OF TRADE RESTRICTION:	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: Restricted to hospital, clinics, and physician offices only: No	Special regulations or returns requirements for this product in certain states?						
Restricted from US territories? (explain in comments) Comments:	If so, which states? Other requirements? Comments?						
MISCELLANIE	OUS NOTES and/or Image of Product Barcode:						
MISCELLANE							



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 S	nip Product	Standard Order Receipt and Processing
Purchase orders may be accepted by: a. EDI		Purchase order daily receipt cut off time by supplier Cut off time:
b. Autofax c. Fax d. Phone only e. Supplier Web Site only Minimum Order Quantity: Supplier's Customer Service Number: Contracted 3PL company / contact #: Phone:	per:	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 Designa	ed Drop Ship Fees:	Overnight and Priority Overnight PO Processing
Expedited freight fees billed with each order: Drop Ship service fee billed with each order:		Overnight receipt available: PO Receipt cut off time:
Drop Ship miscellaneous fees billed: Comments:		Days of week overnight is available: Monday Tuesday Wednesday Thursday Friday
		Priority Overnight receipt available:
Class of Trade Restriction		PO Receipt Cut off time:
No restriction: Select YES if sold to retail pharmacy, hospitals, clinic Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments) Comments:	s and physician offices	Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:
Other Data Information Required to F	rocess 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?
Miscellaneous Notes:		
		ADDITIONAL INFORMATION
		Is product order for scheduled patient procedure? Is product order for restocking purposes?